

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 03:49:50 ; Search time 505.437 Seconds
(without alignments)
8094.223 Million cell updates/sec

Title: US-09-966-264D-1

Perfect score: 137

Sequence: 1 attataaagggaagaaagaaaa.....gtgttgatgtaatt 137

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 27513289 seqs, 14931090276 residues

Total number of hits satisfying chosen parameters: 55026578

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST.*

1: em_estba.*

2: em_esthum.*

3: em_estin.*

4: em_estmu.*

5: em_estov.*

6: em_estpl.*

7: em_estro.*

8: em_htc.*

9: gb_est1.*

10: gb_est2.*

11: gb_hic.*

12: gb_est3.*

13: gb_est4.*

14: gb_est5.*

15: em_estfun.*

16: em_estom.*

17: em_gss_hum.*

18: em_gss_inv.*

19: em_gss_pin.*

20: em_gss_vrt.*

21: em_gss_fun.*

22: em_gss_mam.*

23: em_gss_mus.*

24: em_gss_pro.*

25: em_gss_rod.*

26: em_gss_pig.*

27: em_gss_vrl.*

28: gb_gss1.*

29: gb_gss2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
C 1	137	100.0	373	10 AW580404	AW580404 PM2-HT045
C 2	137	100.0	396	14 H89576	H89576 yw28c12.r1
C 3	137	100.0	455	9 AA427831	AA427831 zw49d12.r
C 4	137	100.0	462	14 CB045405	CB045405 NISC_gc10

C 5	137	100.0	473	14 CA389721	CA389721 cs101f03.
C 6	137	100.0	489	28 AQ015185	AQ015185 CIT-HSP-2
C 7	137	100.0	647	10 AW664684	AW664684 hi84e10.x
C 8	137	100.0	696	10 AW950480	AW950480 EST362550
C 9	137	100.0	710	9 AV725574	AV725574 AV725574
C 10	137	100.0	727	12 BG567176	BG567176 602589640
C 11	137	100.0	740	14 CD356811	CD356811 AGENCOURT
C 12	137	100.0	745	14 CD357556	CD357556 AGENCOURT
C 13	137	100.0	745	14 CD357556	CD357556 AGENCOURT
C 14	137	100.0	1121	12 BM546012	BM546012 AGENCOURT
C 15	136	99.3	743	14 CD110842	CD110842 AGENCOURT
C 16	135.4	98.8	375	10 AW607064	AW607064 PM2-HT045
C 17	135.4	98.8	862	14 CB962272	CB962272 AGENCOURT
C 18	135.4	98.8	911	12 BI752714	BI752714 603028327
C 19	120	87.6	352	10 AW385154	AW385154 PM2-HT045
C 20	111.4	81.3	353	10 AW580423	AW580423 PM2-HT045
C 21	107	78.1	631	14 CB465228	CB465228 726457 MA
C 22	100.6	73.4	251	10 BB339107	BB339107 BB339107
C 23	100.6	73.4	390	9 AI324317	AI324317 mq87c07.y
C 24	100.6	73.4	607	10 BE370292	BE370292 601222549
C 25	100.6	73.4	610	9 AA146038	AA146038 mq87c07.f
C 26	100.6	73.4	735	14 CB570707	CB570707 AGENCOURT
C 27	99	72.3	4437	11 AK036936	AK036936 Mus muscu
C 28	98.6	72.0	296	10 BB041920	BB041920 BB041920
C 29	97.4	71.1	313	10 BB332959	BB332959 BB332959
C 30	95.8	69.9	210	10 BB171938	BB171938 BB171938
C 31	95.8	69.9	268	10 BB178429	BB178429 BB178429
C 32	95.8	69.9	291	10 BB174122	BB174122 BB174122
C 33	95.8	69.9	329	10 BB085328	BB085328 BB085328
C 34	94.6	69.1	841	10 BF791019	BF791019 602251072
C 35	94.2	68.8	293	9 AV307178	AV307178 AV307178
C 36	92.6	67.6	306	10 BB086855	BB086855 BB086855
C 37	92.6	67.6	332	10 BB235445	BB235445 BB235445
C 38	89.6	65.4	418	13 BY377466	BY377466 BY377466
C 39	86.2	62.9	239	9 AV232779	AV232779 AV232779
C 40	86.2	62.9	293	10 BB307434	BB307434 BB307434
C 41	80.8	59.0	641	9 AI528613	AI528613 mq87c07.x
C 42	78.4	57.2	945	10 BF180441	BF180441 601805231
C 43	72	52.6	955	13 BX419179	BX419179 BX419179
C 44	68.4	49.9	651	13 BU338697	BU338697 603515030
C 45	68.4	49.9	733	13 BU257193	BU257193 603744569

ALIGNMENTS

RESULT 1	AW580404	373 bp	mRNA	linear	EST 16-MAR-2000
LOCUS	PM2-HT0451-080100-003-h09	HT0451	Homo sapiens	cdna	mRNA sequence.
DEFINITION	AW580404				
ACCESSION	AW580404.1	GI:7255453			
VERSION	AW580404.1	GI:7255453			
KEYWORDS	EST.				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
REFERENCE	1 (bases 1 to 373)				
AUTHORS	HCGP				
TITLE	The FAPESP/LICR Human Cancer Genome Project				
JOURNAL	Unpublished (1999)				
COMMENT	Contact: Simpson A.J.G. Laboratory of Cancer Genetics Ludwig Institute for Cancer Research Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil Tel: +55-11-2704922 Fax: +55-11-2707001 Email: asimpson@ludwig.org.br This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL (http://www.ludwig.org.br/scripts/gethtml2.pl?tl=PM2&t2=PM2-HT0451-080100-003-h09&l3=2000-01-08&t4=1)				

Seq primer: puc 18 forward
High quality sequence stop: 373.
Location/Qualifiers
1. .373

FEATURES

source

/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_lib="HT0451"
/note="Organ: head neck; Vector: puc18; Site: 1: SmaI;
Site 2: SmaI; A mini-library was made by cloning products
derived from OREGENES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the puc 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."

ORIGIN

Query Match 100.0%; Score 137; DB 10; Length 373;
Best Local Similarity 100.0%; Pred. NO. 4.1e-25;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAAAGAAATAACGCAATGGACAACTGGTGAAGCTGTGAACCTCAGGTGT 60
DB 369 ATTATAAGGAAAAAGAAATAACGCAATGGACAACTGGTGAAGCTGTGAACCTCAGGTGT 310
QY 61 GCACAAATTATCAGGAACACCCCAACCAACCAAGTAGAGTAGAAATAGCATGAGAGCCCGTG 120
DB 309 GCACAAATTATCAGGAACACCCCAACCAACCAAGTAGAGTAGAAATAGCATGAGAGCCCGTG 250
QY 121 TTGTGATGTTAATTAATT 137
DB 249 TTGTGATGTTAATTAATT 233

RESULT 2

H89576/c
LOCUS H89576 396 bp mRNA linear EST 28-NOV-1995
DEFINITION Yw28c12.r1 Morton Fetal Cochlea Homo sapiens cDNA clone
IMAGE:253558 5' similar to gb:M18533 DYSTROPHIN (HUMAN); mRNA
sequence.

ACCESSION H89576.1 GI:1079922

VERSION EST.

KEYWORDS Homo sapiens (human)

SOURCE Homo sapiens

ORGANISM

REFERENCE 1 (bases 1 to 396)

1 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

AUTHORS Hillier, L., Lennon, G., Becker, M., Bonaldo, M.F., Chiapelli, B.,

Chisoe, S., Dietrich, N., Dubuque, T., Favello, A., Gish, W.,

Hawkins, M., Huttman, M., Kucaba, T., Lacy, M., Le, N.,

Mardis, E., Moore, B., Morris, M., Parsons, J., Prange, C., Rifkin, L.,

Rohlfing, T., Scheilenberg, K., Soares, M.B., Tan, F., Thierry-Mieg, J.,

Trevaskis, E., Underwood, K., Wohlmann, P., Waterston, R., Wilson, R.

and Marra, M.

Generation and analysis of 280,000 human expressed sequence tags

Genome Res. 6 (9), 807-828 (1996)

97044478

8889549

Contact: Wilson RK

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: est@watson.wustl.edu

High quality sequence stops: 300

Source: IMAGE Consortium, LLNL

This clone is available royalty-free through LLNL; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.

Insert Length: 1088 Std Error: 0.00

Seq primer: M13Rp1

High quality sequence stop: 300.

FEATURES

source

Location/Qualifiers
1. .396
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="GDB:3891011"
/db_xref="taxon:9606"
/clone="IMAGE:253558"
/tissue_type="cochlea"
/dev_stage="16-22 week fetus"
/lab_host="SOLR cells (Kanamycin resistant)"
/clone_lib="Morton Fetal Cochlea"
/note="Organ: ear; Vector: pBluescript SK-; Site: 1: EcoRI;
Site 2: XhoI; Reference: Genomics 23, 42-50 (1994) Cloned
unidirectionally. Primer: Oligo dt. Fetal cochlea, normal.
37% of inserts <0.5 kb, 56% 0.5-1.0 kb, 7% >1 kb. Uni-ZAP
XR Vector. Library constructed by N. Robertson, C. Morton.
XR adaptor sequence: 5' GAATTCGACGAG 3' ~3' adaptor
sequence: 5' CTCGAGTTTTTTTTTTTTTTT 3' "

ORIGIN

Query Match 100.0%; Score 137; DB 14; Length 396;
Best Local Similarity 100.0%; Pred. No. 4.1e-25;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAAAGAAATAACGCAATGGACAACTGGTGAAGCTGTGAACCTCAGGTGT 60
DB 264 ATTATAAGGAAAAAGAAATAACGCAATGGACAACTGGTGAAGCTGTGAACCTCAGGTGT 205
QY 61 GCACAAATTATCAGGAACACCCCAACCAACCAAGTAGAGTAGAAATAGCATGAGAGCCCGTG 120
DB 204 GCACAAATTATCAGGAACACCCCAACCAACCAAGTAGAGTAGAAATAGCATGAGAGCCCGTG 145
QY 121 TTGTGATGTTAATTAATT 137
DB 144 TTGTGATGTTAATTAATT 128

RESULT 3

AA427831/c

LOCUS

DEFINITION

AA427831 455 bp mRNA linear EST 16-OCT-1997

zw49d12.r1 Soares total fetus Nb2HF8.9w Homo sapiens cDNA clone

IMAGE:773399 5' similar to gb:M18533 DYSTROPHIN (HUMAN); mRNA

sequence.

ACCESSION

AA427831.1 GI:2111628

VERSION EST.

KEYWORDS Homo sapiens (human)

SOURCE Homo sapiens

ORGANISM

REFERENCE

1 (bases 1 to 455)

AUTHORS Hallier, L., Allen, M., Bowles, L., Dubucue, T., Geisel, G., Jost, S.,

Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J.,

Moore, B., Scheilenberg, K., Stepcio, M., Tan, F., Theising, B.,

White, Y., Wylie, T., Waterston, R. and Wilson, R.

WashU-Merck EST Project 1997

Unpublished (1997)

Contact: Wilson RK

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: est@watson.wustl.edu

This clone is available royalty-free through LLNL; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.

Seq primer: -28ml3 rev2 ET from Amersham

High quality sequence stop: 391.

Location/Qualifiers

1. .455

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:773399"

FEATURES

source

constructed by Benito Soares and M. Patricia Boliado.

1	ATTATTAAGGAAAAGAAATATACGCNATGACAAAGTGGTGAAGCTGTGAACTCAGGTGT	60
155	ATTATTAAGGAAAAGAAATATACGCNATGACAAAGTGGTGAAGCTGTGAACTCAGGTGT	96
61	GCACAATTATTCAGGAACACCCCAAAACCAAGTGAAGTAGCAATAGAGAAGCCGCTG	120
95	GCACAATTATTCAGGAACACCCCAAAACCAAGTGAAGTAGCAATAGAGAAGCCGCTG	36
121	TTTGAATGTTAAATTAATT	137
35	TTTGAATGTTAAATTAATT	19

CB045405
 CB045405.1 GI:27783692
 EST.
 Homo sapiens (human)
 Homo sapiens
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 462)
 NC1-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 Unpublished (1997)
 Contact: Robert Strausberg, Ph.D.
 JOURNAL
 COMMENT

DNA Library Preparation: The I.M.A.G.E. Consortium/LLNL
 Sequencing Center (National Institutes of Health Intramural
 Sequencing Center, NCI-CCGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 info@image.llnl.gov
 Plate: i1AM7870 row: E column: 2
 Seq primer: M13RP1 reverse primer (ABI).

```

source
1. "482
location="Guarileis"
organism="Homo sapiens"
mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:3218281"
/tissue_type="juvenile granulosa tumor"
/lab_host="DH10B"
/clone_lib="NCI CGAP Col7"
/notes="Organ: colon; Vector: pCMV-SPORT6; Site 1: SalI;
Site 2: NotI; Cloned unidirectionally. Primer: Oligo dt.
Library constructed by Life Technologies."

```

REFERENCE
AUTHORS
TITLE

JOURNAL
MEDLINE
PUBMED
COMMENT

Mol. Vis. 8 (4), 205-220 (2002)
22103460
12107410

Contact: Wistow G
Section on Molecular Structure and Function
National Eye Institute
6/331 NIH, Bethesda, MD 20892-2740, USA
Tel: 301 402 3452
Fax: 301 496 0078
Email: Graeme@helix.nih.gov

Plate: 101 row: f column: 03
Seq primer: M13RP1 reverse primer (ABI).

```

FEATURES
  source
    seq_params: nixery: reverse primer (50bp).
    Location/Qualifiers
      1..473
        /organism="Homo sapiens"
        /mol_type="mRNA"
        /db_xref="taxon:9606"
        /clone="csl01f03"
        /tissue_type="RPE choroid"
        /dev_stage="Adult"
        /seq_page="EMBL100"

```

```

/seq_node="ENHRL008"
/clone_lib="Human Retinal pigment epithelium/choroid cDNA
(Un-normalized, unamplified): cs"
/notes="Organ: Eye; Vector: pCMVSPORT6; Two different donor
eyes (75-80 years old) yielded approximately 600 mg of
dissected RNA and 7 ug of mRNA. A directionally cloned cDNA
library in the pCMVSPORT6 vector was constructed at Life
Technologies (Rockville, MD; now part of Invitrogen Corp),
essentially following the protocols of the SuperScript
Plasmid System (Invitrogen Corp.
<http://www.invitrogen.com/>). The library code
designation was cs. For this library, cDNA inserts were
cloned into the NotI/Mlu sites of the vector. EST
analysis was performed on the unamplified library at the
NIH Intramural Sequencing Center (NISC)."

```

ORIGIN

Query Match 100.0%; Score 137; DB 14; Length 473;
 Best Local Similarity 100.0%; Pred. No. 3.9e-25;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAATACGCAATGGACAGTGTGAAGCTGTGAACCTCAGGTGT 60
 DB 231 ATTATAAGGAAAAAGAAATACGCAATGGACAGTGTGAAGCTGTGAACCTCAGGTGT 172

QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAATAGCATGAGAACCCGTG 120
 DB 171 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAATAGCATGAGAACCCGTG 112

QY 121 TTGTGATGTTAATTAATT 137
 DB 111 TTGTGATGTTAATTAATT 95

RESULT 6

AQ015185/c

LOCUS AQ015185 489 bp DNA linear GSS 09-JUN-1998
 DEFINITION CIT-HSP-2310F4, TR CIT-HSP Homo sapiens genomic clone 2310F4,
 genomic survey sequence.

ACCESSION AQ015185
 VERSION AQ015185.1 GI:3193921
 KEYWORDS GSS.
 SOURCE Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS Adams,M.D., Rounsley,S.D., Zhao,S., Field,C.E., Bass,S., Linher,K.,
 Golden,K., Berry,K., Granger,D., Suh,E., Wible,C., Shizuya,H.,
 Simon,M. and Venter,J.C.
 TITLE Use of a random BAC End Sequence Database for Sequence-Ready Map
 Building (1998)

JOURNAL

Unpublished (1998)

Contact: Mark Adams

Department of Eukaryotic Genomics

The Institute for Genomic Research

9712 Medical Center Dr., Rockville, MD 20850, USA

Tel: 301 838 0200

Fax: 301 838 0208

Email: mdadams@igrr.org

Clones are available from Research Genetics (info@resgen.com). BAC
 end search page:

http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html.

Seq primer: M13 Reverse

Class: BAC ends.

FEATURES

source

1..489
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /clone="2310F4"
 /sex="Male"
 /cell_type="Sperm"
 /clone_lib="CIT-HSP"
 /notes="Vector: pBelBAC11; Site_1: HindIII; Site_2:
 HindIII"

ORIGIN

Query Match 100.0%; Score 137; DB 28; Length 489;
 Best Local Similarity 100.0%; Pred. No. 3.9e-25;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAATACGCAATGGACAGTGTGAAGCTGTGAACCTCAGGTGT 60
 DB 358 ATTATAAGGAAAAAGAAATACGCAATGGACAGTGTGAAGCTGTGAACCTCAGGTGT 299

QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAATAGCATGAGAACCCGTG 120

ORIGIN

Db 298 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAATAGCATGAGAACCCGTG 239
 QY 121 TTGTGATGTTAATTAATT 137
 Db 238 TTGTGATGTTAATTAATT 222

RESULT 7

AW664684/c

LOCUS AW664684 647 bp mRNA linear EST 06-APR-2000
 DEFINITION h184e10.x1 Soares NFL_T_GBC_S1 Homo sapiens cDNA clone
 IMAGE:2979018 3' similar to gb:M18533 DYSTROPHIN (HUMAN); mRNA
 sequence.

ACCESSION AW664684
 VERSION AW664684.1 GI:7457227
 KEYWORDS EST.
 SOURCE Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 647)

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

AUTHORS National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

TITLE Tumor Gene Index

JOURNAL Unpublished (1997)

COMMENT Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

This clone is available royalty-free through LLNL; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.

Seq primer: -40UP from Gibco

High quality sequence stop: 365.

FEATURES

source

1..647
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:2979018"
 /lab_host="DH10B"
 /clone_lib="Soares NFL_T_GBC_S1"
 /notes="Organ: pooled; Vector: pRT3D-Pac (Pharmacia) with
 a modified polylinker; Site_1: Not 1; Site_2: Eco RI;
 Equal amounts of plasmid DNA from three normalized
 libraries (fetal lung NBHL19W, testis NHT, and B-cell
 NCI CGAP GCBI) were mixed, and ss circles were made in
 vitro. Following HAP purification, this DNA was used as
 tracer in a subtractive hybridization reaction. The driver
 was PCR-amplified cDNAs from pools of 5,000 clones made
 from the same 3 libraries. The pools consisted of
 1.M.A.G.E. clones 297480-302087, 682632-687239,
 726408-728711, and 729096-731399. Subtraction by Bento
 Soares and M. Fatima Bonaldo."

ORIGIN

Query Match 100.0%; Score 137; DB 10; Length 647;
 Best Local Similarity 100.0%; Pred. No. 3.7e-25;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAATACGCAATGGACAGTGTGAAGCTGTGAACCTCAGGTGT 60

Db 151 ATTATAAGGAAAAAGAAATACGCAATGGACAGTGTGAAGCTGTGAACCTCAGGTGT 92

QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAATAGCATGAGAACCCGTG 120

Db 91 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAATAGCATGAGAACCCGTG 32

QY 121 TTGTGATGTTAATTAATT 137

Db 31 TTGTGATGTTAATTAATT 15

RESULT 8

AW950480/c

LOCUS AW950480 696 bp mRNA linear EST 01-JUN-2000
 DEFINITION EST362550 MAGE resequences, MAGA Homo sapiens cDNA, mRNA sequence.
 ACCESSION AW950480
 VERSION AW950480.1 GI:8140134
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 696)
 AUTHORS Hegde, P., Qi, R., Abernathy, K., Dharap, S., Gaspard, R., Gay, C.,
 Holt, I.E., Saeed, A.I., Sharov, V., Lee, N.H., Yeatman, T.J. and
 Quackenbush, J.
 TITLE Assessment of gene expression patterns in a model of colon tumor
 metastasis using a 19,200 element cDNA microarray
 JOURNAL Unpublished (2000)
 COMMENT Contact: John Quackenbush
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 3528
 Fax: 301 838 0208
 Email: johnq@tigr.org
 Plate: 15
 Seq primer: Reverse.
 FEATURES
 source Location/Qualifiers
 1..696
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone_lib="MAGE resequences, MAGA"
 /note="Vector: pBluescriptSKm"
 ORIGIN
 Query Match 100.0%; Score 137; DB 10; Length 696;
 Best Local Similarity 100.0%; Pred. No. 3.7e-25;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ATTATAAGGAAAAAGAAATAACCAATGACCAAGTGTGAAGCTGTGAAGCTGAACTCAGGTGT 60
 DB 572 ATTATAAGGAAAAAGAAATAACCAATGACCAAGTGTGAAGCTGTGAAGCTGAACTCAGGTGT 513
 QY 61 GCACAATTATCAGGAACACCCCAAAACCAAGTGTAGGTAGTAAGTATGATGAGAGCCGCTG 120
 DB 512 GCACAATTATCAGGAACACCCCAAAACCAAGTGTAGGTAGTAAGTATGATGAGAGCCGCTG 453
 QY 121 TTGTGATGTTAATTAATT 137
 DB 452 TTGTGATGTTAATTAATT 436
 RESULT 9
 AV725574/c
 LOCUS AV725574 HTC Homo sapiens cDNA clone HTcBT10 5', mRNA sequence.
 DEFINITION AV725574
 ACCESSION AV725574
 VERSION AV725574.1 GI:10831099
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 710)
 AUTHORS Gu, X., Peng, Y., Song, H., Huang, Q., Yang, Y., Gao, G., Xiao, H., Xu, X.,
 Li, N., Qian, B., Liu, F., Qu, J., Gao, X., Cheng, Z., Xu, Z., Zeng, L.,
 Xu, S., Gu, W., Tu, Y., Jia, J., Fu, G., Ren, S., Zhong, M., Lu, G., Hu, R.,
 Chen, J., Chen, Z. and Han, Z.
 TITLE Homo sapiens cDNA HTC clones
 JOURNAL Unpublished (2000)
 COMMENT Contact: Zeguang Han
 Chinese National Human Genome Center at Shanghai
 351 Guo Shoujing Road, Zhangjiang Hi-Tech Park, Pudong, Shanghai
 201203, P. R. China
 Tel: 86-21-50801919 (ex.45)

Fax: 86-21-50801922
 Email: hanzg@hgc.sh.cn
 This clone is available at CHGC in Shanghai.
 FEATURES
 source Location/Qualifiers
 1..710
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="HTcBT10"
 /tissue_type="Hypothalamus"
 /dev_stage="Adult"
 /lab_host="SOLR"
 /clone_lib="HTC"
 /notes="Vector: pBluescript SK(-); Site_1: EcoRI; Site_2: XhoI"
 ORIGIN
 Query Match 100.0%; Score 137; DB 9; Length 710;
 Best Local Similarity 100.0%; Pred. No. 3.7e-25;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ATTATAAGGAAAAAGAAATAACCAATGACCAAGTGTGAAGCTGTGAAGCTGAACTCAGGTGT 60
 DB 396 ATTATAAGGAAAAAGAAATAACCAATGACCAAGTGTGAAGCTGTGAAGCTGAACTCAGGTGT 337
 QY 61 GCACAATTATCAGGAACACCCCAAAACCAAGTGTAGGTAGTAAGTATGATGAGAGCCGCTG 120
 DB 336 GCACAATTATCAGGAACACCCCAAAACCAAGTGTAGGTAGTAAGTATGATGAGAGCCGCTG 277
 QY 121 TTGTGATGTTAATTAATT 137
 DB 276 TTGTGATGTTAATTAATT 260
 RESULT 10
 BG567176/c
 LOCUS BG567176 727 bp mRNA linear EST 10-APR-2001
 DEFINITION BG567176 NIH_MGC_76 Homo sapiens cDNA clone IMAGE:4723833 5',
 mRNA sequence.
 BG567176
 BG567176.1 GI:13574829
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 727)
 AUTHORS NIH-MGC http://mgc.nci.nih.gov/
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgabbs-r@mail.nih.gov
 Tissue Procurement: CLONETECH Laboratories, Inc.
 cDNA Library Preparation: CLONETECH Laboratories, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov
 Plate: LLCM1583 row: p column: 10
 High quality sequence stop: 694.
 FEATURES
 source Location/Qualifiers
 1..727
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:4723833"
 /lab_host="NIH MGC 76"
 /clone_lib="NIH MGC 76"
 /notes="Organ: liver; Vector: pDNR-LIB (Clontech); Site_1:
 3' adaptors were used in cloning as follows: 5' adaptor
 sequence: 5'-CACGGCCATTATGGCC-3' and 3' adaptor sequence:

Note: this is a NIH MGC Library.

550 ATTATTAAGGAAAAAGAAATACCGAATGGACAGTGGTGAAGCTGTGAACTCAGGTGT 491
Db

61 GCACAAATTATCAGGAACACCCAAAACCAAGTAGAGTAAGCATGAGAACCCGTG 120
QY

490 GCACAAATTATCAGGAACACCCAAAACCAAGTAGAGTAAGCATGAGAACCCGTG 431
Db

121 TTTCATGCTTAATTAATT 137
QY

430 TTTCATGCTTAATTAATT 414
Db

LOCUS	CD357556	745 bp	linear	EST 29-MAY-2003
DEFINITION	AGENCOURT 14253482 NIH MGC 187 Homo sapiens CDNA clone			
	IMAGE:30402369 5', mRNA sequence.			

REFERENCE
1 (bases 1 to 745)
AUTHORS
NIH-MGC <http://mgc.nci.nih.gov/>.
TITLE
National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL
Unpublished (1999)
CONTACT
Contact: Daniela S. Gerhard, Ph.D.
COMMENT

DNA sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
Plate: NDCM184 row: j column: 10
High quality sequence stop: 460.

```

/clones="IMAGE:30402369"
/lab_host="DH10B (T1 phage-resistant)"
/clone_lib="NIH_MGC_187"

note="Organ: Blood vessels - aorta, basilar and artery;
Vector: pDNR-LIB; Site_1: SfiI (ggccattatggcc); Site_2:
SfiI (ggccgctcgccc); 5' and 3' adaptors were used in
cloning as follows: 5' adaptor sequence:
5'-CACGCCCATATGCCC-3' and 3' adaptor sequence:
5'-ATTCTAGAGCCCGGCGCCGACATG-dT(30)BN-3' (where B = A,
C, or G and N = A, C, G, or T). Average insert size 1.4 kb
(range 0.5-4.0 kb). 14/15 colonies contained inserts by
PCR. This library was enriched for full-length clones and
was constructed by Clontech Laboratories (Palo Alto, CA).
Note: this is a NIH MGC library."

```

Query Match 100.0%; Score 137; DB 14; Length 745;
Best Local Similarity 100.0%; Pred. No. 3.6e-25;

[illegible]

Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAAGAAATAACCAATGCAAGTGGTGAAGCTGTGAACCTCAGGTGT 60
 DB 260 ATTATAAGGAAAAAAGAAATAACCAATGCAAGTGGTGAAGCTGTGAACCTCAGGTGT 201

QY 61 GCACAATTATCAGGAACACCCCAACCAACCAAGTGGTAGAAATAGCATGAGAGCCGTG 120
 DB 200 GCACAATTATCAGGAACACCCCAACCAACCAAGTGGTAGAAATAGCATGAGAGCCGTG 141

QY 121 TTGTGATGTTAATTAATT 137
 DB 140 TTGTGATGTTAATTAATT 124

RESULT 13
 BM546012/c
 LOCUS
 DEFINITION BM546012 1121 bp mRNA linear EST 20-FEB-2002
 AGENCOURT_6505286 NIH_MGC_125 Homo sapiens cDNA clone IMAGE:5588419
 5', mRNA sequence.

ACCESSION BM546012
 VERSION BM546012.1 GI:18778623
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 1121)
 AUTHORS NIH-MGC http://mgi.nci.nih.gov/
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: Invitrogen
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone Distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
 Plate: LLNL2359 row: h column: 20
 High quality sequence start: 50
 High quality sequence stop: 764.

FEATURES
 Location/Qualifiers
 1..1121
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:5588419"
 /lab_host="DH10B"
 /clone_lib="NIH MGC 125"
 /note="Organ: ovary (pool of 3); Vector: pCMV-SPORT6; Site 1: EcoRV (destroyed); Site 2: NotI; RNA source pool of three ovaries, from females ranging in age from 38 to 49 yo. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 2.1 kb, insert size range 1-3.5 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 036."

ORIGIN
 Query Match 100.0%; Score 137; DB 12; Length 1121;
 Best Local Similarity 100.0%; Pred. No. 3.4e-25;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAAGAAATAACGCAATGGCAAGTGGTGAAGCTGTGAACCTCAGGTGT 60
 DB 514 ATTATAAGGAAAAAAGAAATAACGCAATGGCAAGTGGTGAAGCTGTGAACCTCAGGTGT 455

QY 61 GCACAATTATCAGGAACACCCCAACCAACCAAGTGGTAGAAATAGCATGAGAGCCGTG 120
 DB 454 GCACAATTATCAGGAACACCCCAACCAACCAAGTGGTAGAAATAGCATGAGAGCCGTG 395

QY 121 TTGTGATGTTAATTAATT 137
 DB 394 TTGTGATGTTAATTAATT 378

RESULT 14
 CD110642/c
 LOCUS
 DEFINITION CD110642 743 bp mRNA linear EST 15-MAY-2003
 AGENCOURT_13995397 NIH_MGC_187 Homo sapiens cDNA clone IMAGE:30373580 5', mRNA sequence.

ACCESSION CD110642
 VERSION CD110642.1 GI:30754851
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 743)
 AUTHORS NIH-MGC http://mgi.nci.nih.gov/
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: Dr. Michael Brownstein and Dr. Miklos Palkovits
 cDNA Library Preparation: CLONTECH Laboratories, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone Distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
 Plate: NDCM167 row: j column: 21
 High quality sequence stop: 578.

FEATURES
 Location/Qualifiers
 1..743
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:30373580"
 /lab_host="DH10B (T1 phage-resistant)"
 /clone_lib="NIH MGC 187"
 /note="Organ: Blood_vessels - aorta, basilar and artery; Vector: pMNR-LIB; Site 1: SfiI (ggccattatggcc); Site 2: SfiI (ggccgctcgcc); 5' and 3' adaptors were used in cloning as follows: 5' adaptor sequence: 5'-CACGCCATTATGCCC-3' and 3' adaptor sequence: 5'-ATTCTAGAGCCGAGCGCGCCGACATG-dt(30)BN-3' (where B = A, C, or G and N = A, C, G, or T). Average insert size 1.4 kb (range 0.5-4.0 kb). 14/15 colonies contained inserts by PCR. This library was enriched for full-length clones and was constructed by Clontech Laboratories (Palo Alto, CA). Note: this is a NIH_MGC Library."

ORIGIN
 Query Match 99.3%; Score 136; DB 14; Length 743;
 Best Local Similarity 100.0%; Pred. No. 6.6e-25;
 Matches 136; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAAGAAATAACGCAATGGCAAGTGGTGAAGCTGTGAACCTCAGGTGT 60
 DB 138 ATTATAAGGAAAAAAGAAATAACGCAATGGCAAGTGGTGAAGCTGTGAACCTCAGGTGT 79

QY 61 GCACAATTATCAGGAACACCCCAACCAACCAAGTGGTAGAAATAGCATGAGAGCCGTG 120
 DB 78 GCACAATTATCAGGAACACCCCAACCAACCAAGTGGTAGAAATAGCATGAGAGCCGTG 19

QY 121 TTGTGATGTTAATTAATT 136
 DB 18 TTGTGATGTTAATTAATT 3

RESULT 15
 AW607064/c

LOCUS AW607064 375 bp mRNA linear EST 23-MAR-2000
 DEFINITION PM2-HT0451-170100-004-h12 HT0451 Homo sapiens cDNA, mRNA sequence.
 ACCESSION AW607064
 VERSION AW607064.1 GI:7311805
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 375)
 HCGP <http://www.ludwig.org.br/ORESTES>.
 The FAPESP/LICR Human Cancer Genome Project
 Unpublished (1999)
 CONTACT: Simpson A.J.G.
 Laboratory of Cancer Genetics
 Ludwig Institute for Cancer Research
 Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
 Brazil
 Tel: +55-11-2704922
 Fax: +55-11-2707001
 Email: asimpson@ludwig.org.br
 This sequence was derived from the FAPESP/LICR Human Cancer Genome
 Project. This entry can be seen in the following URL
 (<http://www.ludwig.org.br/scripts/gethtml2.pl?t1=PM2&t2=PM2-HT0451-170100-004-h12&t3=2000-01-17&t4=1>)
 Seq primer: puc 18 forward
 High quality sequence stop: 375.
 Location/Qualifiers
 1..375
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /dev_stage="Adult"
 /clone_lib="HT0451"
 /note="Organ: head neck; Vector: puc18; Site 1: Smal;
 Site 2: Smal; A mini-library was made by cloning products
 derived from ORESTES PCR (U.S. Letters Patent application
 No. 196,716 - Ludwig Institute for Cancer Research)
 profiles into the pUC 18 vector. Reverse transcription of
 tissue mRNA and cDNA amplification were performed under
 low stringency conditions."
 ORIGIN
 Query Match 98.8%; Score 135.4; DB 10; Length 375;
 Best Local Similarity 99.3%; Pred. No. 1.1e-24;
 Matches 136; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 ATTATAAGGAAAAAGAAATAACGCAATGGACAGCTGTGAAGCTGTGAACCTCAGGTGT 60
 ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
 Db 371 ATTATAAGGAAAAAGAAATAACGCAATGGACAGCTGTGAAGCTGTGAACCTCAGGTGT 312
 ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
 QY 61 GCACATATTCAGGAACACCCCAACCAAGTGGTAGAATAATAGCATGAGAGCCGCG 120
 ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
 Db 311 GCACATATTCAGGAACACCCCAACCAAGTGGTAGAATAATAGCATGAGAGCCGCG 252
 ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
 QY 121 TTGATGTTAATTAAT 137
 ||||||||||||||||
 Db 251 TTGATGTTAATTAAT 235
 ||||||||||||||||

Search completed: April 6, 2004, 14:12:02
 Job time : 511.437 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 02:56:20 ; Search time 64.812 Seconds
(without alignments)

8979.866 Million cell updates/sec

Title: US-09-966-264D-1

Perfect score: 137

Sequence: 1 attataaaggaaagaaaa.....gtgttgatgtaattaatt 137

Scoring table: IDENTITY NUC

Gapop 10_0 , Gapext 1.0

Searched: 337863 seqs, 2124099041 residues

Total number of hits satisfying chosen parameters: 6747726

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N Geneseq 29Jan04:*

- 1: Geneseqn1980a:*
- 2: Geneseqn1990a:*
- 3: Geneseqn1990b:*
- 4: Geneseqn2000a:*
- 5: Geneseqn2001a:*
- 6: Geneseqn2002a:*
- 7: Geneseqn2003a:*
- 8: Geneseqn2003bs:*
- 9: Geneseqn2003cs:*
- 10: Geneseqn2004a:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	137	100.0	137	6	ABK86496 Human Apo
C 2	137	100.0	158	6	ABK86469 Human Apo
C 3	137	100.0	200	6	ABK86468 Human Apo
C 4	137	100.0	455	9	ADD32504 Human mit
5	137	100.0	996	6	ABK86497 Human Apo
6	137	100.0	1230	6	ABK86462 Human Apo
7	137	100.0	1234	6	ABK86463 Human Apo
C 8	137	100.0	2691	6	ABK81996 Human Apo
C 9	137	100.0	8689	6	ABK82000 DNA encod
C 10	137	100.0	11443	6	ABK82002 DNA encod
C 11	137	100.0	13957	6	ABK81959 CDNA enco
C 12	137	100.0	13957	6	ABT10904 Human bre
C 13	137	100.0	13957	6	ABN95786 Gene #228
C 14	137	100.0	13957	6	ABK86900 Human dys
C 15	126	92.0	13977	6	ABK870403 Human bon
C 16	107	78.1	959	6	ABN74601 Bovine em
C 17	100.6	73.4	13815	2	ABN18885 Mus muscu
C 18	100.6	73.4	13815	6	ABK81960 CDNA enco
C 19	100.6	73.4	13815	6	ABN99799 Mouse isc
C 20	100.6	73.4	19307	2	AAT27558 Shuttle v
C 21	57.2	41.8	108	6	ABK86467 Human apo
C 22	41	29.9	60	6	ABK86471 Human apo
C 23	33.6	24.5	1863	9	ADK60952 Human gen

C 24	31.8	23.2	6428	4	AAK78562	Human imm
C 25	31.8	23.2	96591	9	ADC85301	Mouse Sos
C 26	31.8	23.2	96592	8	ADA02822	Human SOS
C 27	31.8	23.2	96592	9	ADB72560	Human SOS
C 28	31.6	23.1	2000	6	ABZ17261	Arabidops
C 29	31.2	22.8	1977	6	ABL62657	Colion ace
C 30	31.2	22.8	1977	6	ABT10871	Human bre
C 31	31.2	22.8	1977	6	ABK83769	Human CDN
C 32	31.2	22.8	1979	2	AAT87015	Human FC-
C 33	31.2	22.8	1990	1	AAN92451	cDNA inse
C 34	31.2	22.8	2461	4	AAS94845	Human DNA
C 35	30.8	22.5	47319	4	AAK72230	Human imm
C 36	30.8	22.5	47319	4	AAK64813	Human imm
C 37	30.8	22.5	236303	4	AAS11614	Human gen
C 38	30.6	22.3	538	6	ABL78454	Human Ova
C 39	30.6	22.3	539	6	ABK62246	Rat sequ
C 40	30.6	22.3	539	7	ABT40428	Toxicity
C 41	30.6	22.3	539	9	ADB55325	Toxicity
C 42	30.6	22.3	539	9	ADB49833	Primary r
C 43	30.6	22.3	6644	2	AAK33181	Base sequ
C 44	30.6	22.3	7372	2	AAK33182	Base sequ
C 45	30.6	22.3	7797	2	AAK33180	Cowpox vi

ALIGNMENTS

RESULT 1
ABK86496
ID ABK86496 standard; DNA; 137 BP.
XX
AC ABK86496;
XX
DT 27-AUG-2002 (first entry)
XX
DE Human Apo-dystrophin-4 inversion sequence.
XX
KW Human; ds; apo-dystrophin-4; inversion sequence; gene therapy;
KW protein truncation; muscular dystrophy; leukaemia.
XX
OS Homo sapiens.
XX
FH Key
FT CDS
FT
FT Location/Qualifiers
3. .137
/*tag= a
/product= "apo-dystrophin-4 peptide appearing as
AAU98738"
/partial
/note= "No start or stop codon shown"
/transl_except= (pos:21..23,aa:Xaa)
/transl_except= (pos:48..50,aa:Xaa)
/transl_except= (pos:93..95,aa:Xaa)
/transl_except= (pos:123..125,aa:Xaa)
/transl_except= (pos:129..131,aa:Xaa)
/note= "Xaa= unknown, encoded by in frame stop codon"

GB2368064-A.

24-APR-2002.

16-JAN-2001; 2001GB-00001124.

30-SEP-2000; 2000US-0237079P.

(IMCR) IMPERIAL CANCER RES TECHNOLOGY LTD.

(BARB/) BARBER E.

Barber E;

WPI; 2002-429042/46.

P-PSDB; AAU98737.

PT New human regulatory polynucleotide, useful for treating disorders

PT associated with protein truncation, particularly muscular dystrophy, and
 PT related peptides and antibodies.
 XX Claim 1; Page 169; 222pp; English.

XX The invention relates to a polynucleotide (I) comprising, or consisting
 CC of, apo-dystrophin-4 inversion sequence appearing as ABK86496, or its
 CC functional equivalents (e.g. the apo-dystrophin-4 cDNA sequence appearing
 CC as ABK86497). Also included are polynucleotides that hybridize to either
 CC strand of (I), a vector containing (I), a cell containing (I) or the
 CC vector, proteins and peptides encoded by (I), a protein homologous with
 CC human dystrophin that is expressed on cell surfaces in vivo antibodies
 CC (Ab) specific for the protein and method of screening for leukemia cells
 CC The apo-dystrophin-4 inversion sequence is a regulatory element that
 CC controls expression (transcription and translation) of associated DNA,
 CC and may allow read-through of stop codons. The apo-dystrophin-4 inversion
 CC sequence is used in gene therapy of diseases associated with truncation
 CC of proteins, particularly muscular dystrophy and also leukaemia, but more
 CC generally (I) is a regulatory sequence used to control expression of any
 CC attached gene. Analysis of DNA for (I), or detection of proteins (II)
 CC encoded by (I), can be used to screen for leukaemic cells and related
 CC diseases. Antibodies raised against (II) can be used therapeutically, to
 CC inhibit (II) activity, also to detect (II) in screening assays. The
 CC present sequence is the apo-dystrophin-4 inversion sequence and upstream
 CC genomic region

XX Sequence 137 BP; 56 A; 19 C; 32 G; 30 T; 0 U; 0 Other;

Query Match 100.0%; Score 137; DB 6; Length 137;
 Best Local Similarity 100.0%; Pred. No. 1.6e-33;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAATAACGCAATGGACAGTGGTGAAGCTGTGAAGTGTGAGTGTG 60

DB 1 ATTATAAGGAAAAAGAAATAACGCAATGGACAGTGGTGAAGCTGTGAGTGTGAGTGTG 60

QY 61 GCACATTATCAGGAACACCCCAACCAAGTGGTGAAGTGTGAAGTGTGAAGTGTGAAGTGTG 120

DB 61 GCACATTATCAGGAACACCCCAACCAAGTGGTGAAGTGTGAAGTGTGAAGTGTGAAGTGTG 120

QY 121 TTGTGTTTAATTAATT 137

DB 121 TTGTGTTTAATTAATT 137

RESULT 2

ABK86469/C
 ID ABK86469 standard; cDNA; 158 BP.

XX ABK86469;

XX 27-AUG-2002 (first entry)

XX Human apo-dystrophin-4 cDNA fragment with inversion breakpoint #2.

XX Human; ss; apo-dystrophin-4; inversion sequence; gene therapy;
 KW protein truncation; muscular dystrophy; leukaemia; dystrophin.

XX Homo sapiens.

XX Key Location/Qualifiers
 FT misc_recomb 137
 FT /*tag= a
 FT /label= inversion_breakpoint

XX CB2368064-A.

XX 24-APR-2002.

XX 16-JAN-2001; 2001GB-00001124.

XX 30-SEP-2000; 2000US-0237079P.

XX (IMCR) IMPERIAL CANCER RES TECHNOLOGY LTD.
 PA (BARB/) BARBER E.

XX Barber E;

XX WPI; 2002-429042/46.

XX New human regulatory polynucleotide, useful for treating disorders
 PT associated with protein truncation, particularly muscular dystrophy, and
 PT related peptides and antibodies.

XX Disclosure; Fig 16B; 222pp; English.

XX The invention relates to a polynucleotide (I) comprising, or consisting
 CC of, apo-dystrophin-4 inversion sequence appearing as ABK86496, or its
 CC functional equivalents (e.g. the apo-dystrophin-4 cDNA sequence appearing
 CC as ABK86497). Also included are polynucleotides that hybridize to either
 CC strand of (I), a vector containing (I), a cell containing (I) or the
 CC vector, proteins and peptides encoded by (I), a protein homologous with
 CC human dystrophin that is expressed on cell surfaces in vivo antibodies
 CC (Ab) specific for the protein and method of screening for leukemia cells
 CC by analysing DNA for presence of (I) or by detecting presence of (II).
 CC The apo-dystrophin-4 inversion sequence is a regulatory element that
 CC controls expression (transcription and translation) of associated DNA,
 CC and may allow read-through of stop codons. The apo-dystrophin-4 inversion
 CC sequence is used in gene therapy of diseases associated with truncation
 CC of proteins, particularly muscular dystrophy and also leukaemia, but more
 CC generally (I) is a regulatory sequence used to control expression of any
 CC attached gene. Analysis of DNA for (I), or detection of proteins (II)
 CC encoded by (I), can be used to screen for leukaemic cells and related
 CC diseases. Antibodies raised against (II) can be used therapeutically, to
 CC inhibit (II) activity, also to detect (II) in screening assays. The
 CC present sequence is a apo-dystrophin-4 cDNA fragment showing an inversion
 CC breakpoint (recombination signal sequence) similar to that in with human
 CC dystrophin

XX Sequence 158 BP; 35 A; 35 C; 20 G; 68 T; 0 U; 0 Other;

Query Match 100.0%; Score 137; DB 6; Length 158;
 Best Local Similarity 100.0%; Pred. No. 1.7e-33;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAATAACGCAATGGACAGTGGTGAAGCTGTGAAGTGTGAGTGTG 60

DB 137 ATTATAAGGAAAAAGAAATAACGCAATGGACAGTGGTGAAGCTGTGAAGTGTGAGTGTG 78

QY 61 GCACATTATCAGGAACACCCCAACCAAGTGGTGAAGTGTGAAGTGTGAAGTGTGAAGTGTG 120

DB 77 GCACATTATCAGGAACACCCCAACCAAGTGGTGAAGTGTGAAGTGTGAAGTGTGAAGTGTG 120

QY 121 TTGTGTTTAATTAATT 137

DB 17 TTGTGTTTAATTAATT 1

RESULT 3

ABK86468/C

ID ABK86468 standard; DNA; 200 BP.

XX ABK86468;

XX 27-AUG-2002 (first entry)

XX Human dystrophin genomic DNA with inversion breakpoint #2.

XX Human; ds; apo-dystrophin-4; inversion sequence; gene therapy;
 KW protein truncation; muscular dystrophy; leukaemia; dystrophin.

XX Homo sapiens.

XX Key Location/Qualifiers
 FT misc_recomb 42

```

FT FT      /tag= a
FT FT      /label= Inversion_breakpoint
FT FT      179
FT FT      /tag= b
FT FT      /label= Inversion_breakpoint
XX
PN GB2368064-A.
PD 24-APR-2002.
XX
PF 16-JAN-2001; 2001GB-00001124.
XX
PR 30-SEP-2000; 2000US-0237079P.
XX
PA (IMCR ) IMPERIAL CANCER RES TECHNOLOGY LTD.
PA (BARB/) BARBER E.
XX
PI Barber E;
XX
XX WPI; 2002-429042/46.
DR
XX New human regulatory polynucleotide, useful for treating disorders
PT associated with protein truncation, particularly muscular dystrophy, and
PT related peptides and antibodies.
XX
PS Disclosure; Fig 16B; 222pp; English.
XX
CC The invention relates to a polynucleotide (I) comprising, or consisting
CC of, apo-dystrophin-4 inversion sequence appearing as ABK86496, or its
CC functional equivalents (e.g. the apo-dystrophin-4 cDNA sequence appearing
CC as ABK86497). Also included are polynucleotides that hybridise to either
CC strand of (I), a vector containing (I), a cell containing (I) or the
CC vector, proteins and peptides encoded by (I), a protein homologous with
CC human dystrophin that is expressed on cell surfaces in vivo antibodies
CC (Ab) specific for the protein and method of screening for leukemia cells
CC by analysing DNA for presence of (I) or by detecting presence of (II).
CC The apo-dystrophin-4 inversion sequence is a regulatory element that
CC controls expression (transcription and translation) of associated DNA,
CC and may allow read-through of stop codons. The apo-dystrophin-4 inversion
CC sequence is used in gene therapy of diseases associated with truncation
CC of proteins, particularly muscular dystrophy and also leukaemia, but more
CC generally (i) is a regulatory sequence used to control expression of any
CC attached gene. Analysis of DNA for (I), or detection of proteins (II)
CC encoded by (I), can be used to screen for leukaemic cells and related
CC diseases. Antibodies raised against (II) can be used therapeutically, to
CC inhibit (II) activity, also to detect (II) in screening assays. The
CC present sequence is a human dystrophin genomic DNA fragment showing
CC inversion breakpoints (recombination signal sequence) similar to that in
CC apo-dystrophin-4
XX
SQ Sequence 200 BP; 44 A; 40 C; 28 G; 88 T; 0 U; 0 Other;
Query Match 100.0%; Score 137; DB 6; Length 200;
Best Local Similarity 100.0%; Pred. No. 1.8e-33;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAAAGAAATAACGCAATGGCAAGTGTGTGAAGCTGTGAACCTCAGGTGT 60
Db 179 ATTATAAGGAAAAAGAAATAACGCAATGGCAAGTGTGTGAAGCTGTGAACCTCAGGTGT 120
QY 61 GCACATTTATCAGGACACCCCAACCAACCAAGTGTAGTAAATAGCATGAGACCCGTG 120
Db 119 GCACATTTATCAGGACACCCCAACCAACCAAGTGTAGTAAATAGCATGAGACCCGTG 60
QY 121 TTGTATGTTTAATTAATT 137
Db 59 TTGTATGTTTAATTAATT 43
XX
RESULT 4
ADD32504/c
ID ADD32504 standard; DNA; 455 BP.
XX

```

```

AC ADD32504;
XX
XX 15-JAN-2004 (first entry)
XX
DE Human mitochondrial DNA sequence SEQ ID NO:274.
XX
XX ds; human; array; mitochondrial; hybridisation; energy-metabolism;
XX mitochondrial disease; oxidative phosphorylation dysfunction;
XX oxidative stress; apoptosis; aging.
XX
XX Homo sapiens.
XX
XX WO2003020220-A2.
XX
XX 13-MAR-2003.
XX
XX 30-AUG-2002; 2002WO-US027886.
XX
XX 30-AUG-2001; 2001US-0316323P.
XX
XX 31-AUG-2001; 2001CA-02356540.
XX
XX (UYEM-) UNIV EMORY.
XX
XX Wallace DC, Levy S, Kerstann K, Procaccio V;
XX
XX WPI; 2003-300821/29.
XX
XX Array containing probes for genes involved in mitochondrial biology,
XX useful for determining mitochondrial biology gene expression profiles for
XX use in diagnosing pathologies and identifying biochemical pathways.
XX
XX Claim 2; SEQ ID NO 274; 201pp; English.
XX
XX The invention relates to a novel array comprising at least two isolated
XX nucleotide molecules, each molecule having a sequence capable of uniquely
XX hybridising to a nucleic acid molecule which is an expression product of
XX a gene involved in mitochondrial biology. The array comprises two or more
XX isolated nucleic acid molecules or spots, each molecule having a sequence
XX chosen from sequence of 394 human probes and 2046 mouse probes. An array
XX of the invention is useful for determining an expression profile of a
XX mouse or human sample containing nucleic acid, by contacting the array
XX with the sample under conditions allowing selective hybridisation, and
XX measuring hybridisation of nucleic acid in the sample to the array to
XX produce an expression profile. The array is also useful for determining
XX an expression profile of a first labelled sample containing nucleic acid
XX relative to a second, differently labelled sample containing nucleic
XX acid. The second sample is a reference or a standard. An array is useful
XX for determining an expression profile diagnostic of an energy-metabolism-
XX related physiological condition. An array of the invention is useful for
XX determining mitochondrial biology gene expression profiles of organisms,
XX such as human, mice and closely related species, tissue and organs of
XX such organisms, which are useful for determining expression profiles
XX diagnostic of energy metabolism-related physiological conditions
XX diagnosing such physiological conditions, identifying biochemical
XX pathways, genes, and mutations involved in such physiological conditions,
XX identifying therapeutic agents useful for preventing and/or treating such
XX physiological conditions, evaluating and/or monitoring the efficacy of
XX such therapies, and creating and identifying animal models of human
XX energy metabolism-related physiological conditions. An array is also
XX useful for defining expression signatures or profiles for mitochondrial
XX diseases, as well as distinguishing clinical disorders that result from
XX oxidative phosphorylation (OXPHOS) dysfunction, oxidative stress,
XX apoptosis and aging. An array of the invention contains probes of genes
XX not previously recognised to participate in mitochondrial biology. The
XX sequences shown in ADD3231-ADD33223 represent human mitochondrial DNA
XX clones used to make the probes of the invention. Some sequences are not
XX present, these are SEQ ID NO's 295, 1174, 1213, 1700, 1728, 1730, 1905,
XX 1906, 2408 and 2643.
XX
XX Sequence 455 BP; 127 A; 80 C; 63 G; 185 T; 0 U; 0 Other;
Query Match 100.0%; Score 137; DB 9; Length 455;
Best Local Similarity 100.0%; Pred. No. 2.3e-33;

```

```
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAAAGAAATAACGCAATGGACAAAGTGGTGAAGCTGGAAGTCAAGTGT 60
Dd 155 ATTATAAGGAAAAAGAAATAACGCAATGGACAAAGTGGTGAAGCTGGAAGTCAAGTGT 96
QY 61 GCACAAATTATCAGGAGACACCCCAACCAACCAAGTGGTGAAGTGAAGTGAAGTGAAGT 120
Dd 95 GCACAAATTATCAGGAGACACCCCAACCAACCAAGTGGTGAAGTGAAGTGAAGTGAAGT 36
QY 121 TTGTGATTTAATTAATT 137
Dd 35 TTGTGATTTAATTAATT 19
RESULT 5
ABK86497
ID ABK86497 standard; cDNA; 996 BP.
AC ABK86497;
XX
XX
DT 27-AUG-2002 (first entry)
XX
DE Human Apo-dystrophin-4 cDNA.
XX
KW Human; ss; gene; apo-dystrophin-4; inversion sequence; gene therapy;
KW protein truncation; muscular dystrophy; leukaemia.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT CDS 1..996
FT /tag= a
FT /product= "Apo-dystrophin-4 protein appearing as
AAU98739"
FT /partial
FT /note= "No start or stop codon shown"
FT /transl_except= (pos:7..9,aa:Xaa)
FT /transl_except= (pos:49..51,aa:Xaa)
FT /transl_except= (pos:187..189,aa:Xaa)
FT /transl_except= (pos:202..204,aa:Xaa)
FT /transl_except= (pos:217..219,aa:Xaa)
FT /transl_except= (pos:253..255,aa:Xaa)
FT /transl_except= (pos:265..270,aa:Xaa-Xaa)
FT /transl_except= (pos:394..396,aa:Xaa)
FT /transl_except= (pos:547..549,aa:Xaa)
FT /transl_except= (pos:565..567,aa:Xaa)
FT /transl_except= (pos:616..618,aa:Xaa)
FT /transl_except= (pos:649..651,aa:Xaa)
FT /transl_except= (pos:682..684,aa:Xaa)
FT /transl_except= (pos:709..711,aa:Xaa)
FT /transl_except= (pos:730..732,aa:Xaa)
FT /transl_except= (pos:841..843,aa:Xaa)
FT /transl_except= (pos:850..852,aa:Xaa)
FT /transl_except= (pos:880..882,aa:Xaa)
FT /transl_except= (pos:907..909,aa:Xaa)
FT /transl_except= (pos:952..954,aa:Xaa)
FT /transl_except= (pos:982..984,aa:Xaa)
FT /transl_except= (pos:988..990,aa:Xaa)
FT /note= "Xaa= unknown, encoded by in frame stop codon"
XX
XX
PN GB2368064-A.
XX
XX
PD 24-APR-2002.
XX
XX
PF 16-JAN-2001; 2001GB-00001124.
XX
XX
PR 30-SEP-2000; 2000US-0237079P.
XX
XX
PA (IMCR ) IMPERIAL CANCER RES TECHNOLOGY LTD.
PA (BARB/) BARBER E.
XX
PI Barber E;
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```
XX WPI; 2002-429042/46.
DR P-PSDB; AAU98739.
XX
XX New human regulatory polynucleotide, useful for treating disorders
PT associated with protein truncation, particularly muscular dystrophy, and
PT related peptides and antibodies.
XX
XX Claim 5; Page 170-172; 222pp; English.
XX
CC The invention relates to a polynucleotide (I) comprising, or consisting
CC of, apo-dystrophin-4 inversion sequence appearing as ABK86496, or its
CC functional equivalents (e.g. the apo-dystrophin-4 cDNA sequence appearing
CC as ABK86497). Also included are polynucleotides that hybridize to either
CC strand of (I), a vector containing (I), a cell containing (I) or the
CC vector, proteins and peptides encoded by (I), a protein homologous with
CC human dystrophin that is expressed on cell surfaces in vivo antibodies
CC (Ab) specific for the protein and method of screening for leukemia cells
CC by analysing DNA for presence of (I) or by detecting presence of (II).
CC The apo-dystrophin-4 inversion sequence is a regulatory element that
CC controls expression (transcription and translation) of associated DNA,
CC and may allow read-through of stop codons. The apo-dystrophin-4 inversion
CC sequence is used in gene therapy of diseases associated with truncation
CC of proteins, particularly muscular dystrophy and also leukaemia, but more
CC generally (I) is a regulatory sequence used to control expression of any
CC attached gene. Analysis of DNA for (I), or detection of proteins (II)
CC encoded by (I), can be used to screen for leukaemic cells and related
CC diseases. Antibodies raised against (II) can be used therapeutically, to
CC inhibit (II) activity, also to detect (II) in screening assays. The
CC present sequence is the cDNA sequence for human apo-dystrophin-4
CC containing a plurality of stop codons some of which may be read through
CC due to the presence of (I) in the apo-dystrophin-4 gene
XX
XX Sequence 996 BP; 334 A; 152 C; 195 G; 315 T; 0 U; 0 Other;
Query Match 100.0%; Score 137; DB 6; Length 996;
Best Local Similarity 100.0%; Pred. No. 2.8e-33;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAAAGAAATAACGCAATGGACAAAGTGGTGAAGCTGGAAGTCAAGTGT 60
Dd 860 ATTATAAGGAAAAAGAAATAACGCAATGGACAAAGTGGTGAAGCTGGAAGTCAAGTGT 919
QY 61 GCACAAATTATCAGGAGACACCCCAACCAACCAAGTGGTGAAGTGAAGTGAAGTGAAGT 120
Dd 920 GCACAAATTATCAGGAGACACCCCAACCAACCAAGTGGTGAAGTGAAGTGAAGTGAAGT 979
QY 121 TTGTGATTTAATTAATT 137
Dd 980 TTGTGATTTAATTAATT 996
RESULT 6
ABK86462
ID ABK86462 standard; cDNA; 1230 BP.
XX
XX AC ABK86462;
XX
XX DT 27-AUG-2002 (first entry)
XX
XX DE Human Apo-dystrophin-4 full length cDNA.
XX
XX KW Human; ss; gene; apo-dystrophin-4; inversion sequence; gene therapy;
KW protein truncation; muscular dystrophy; leukaemia.
XX
XX OS Homo sapiens.
XX
XX FH Key Location/Qualifiers
XX FT CDS 1..1230
XX FT /tag= a
XX FT /product= "Apo-dystrophin-4"
XX FT /partial
XX FT /note= "No start or stop codon shown"
```


XX SQ Sequence 1234 BP; 405 A; 190 C; 260 G; 379 T; 0 U; 0 Other;
 Query Match 100.0%; Score 137; DB 6; Length 1234;
 Best Local Similarity 100.0%; Pred. No. 3e-33;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAATAACGCAATGGCAAGTGGTAGAAGCTGTGAAGCTCAGGTGT 60
 DB 1098 ATTATAAGGAAAAAATAACGCAATGGCAAGTGGTAGAAGCTGTGAAGCTCAGGTGT 1157

QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAAGCTGTGAAGCTCAGGTGT 120
 DB 1158 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAAGCTGTGAAGCTCAGGTGT 1217

QY 121 TTGTGTTTAATTAATT 137
 DB 1218 TTGTGTTTAATTAATT 1234

RESULT 8
 ABK81996/c
 ID ABK81996 standard; DNA; 2691 BP.
 XX AC ABK81996;
 XX DT 13-AUG-2002 (first entry)
 XX DE Human dystrophin 3' untranslated region.
 XX KW Mini-dystrophin peptide; spectrin-like repeat domain; muscle disease;
 XX KW Duchenne's muscular dystrophy; DMD; dystrophin; human; ds.
 XX OS Homo sapiens.
 XX DN WO200229056-A2.
 XX PD 11-APR-2002.
 XX PF 04-OCT-2001; 2001WO-US0311126.
 XX PR 06-OCT-2000; 2000US-0238848P.
 XX PA (UNMI) UNIV MICHIGAN.
 XX PI Chamberlain JS, Harper SQ;
 XX DR WPI; 2002-435334/46.
 XX CC The invention describes a composition comprising a mini-dystrophin peptide comprising a spectrin-like repeat domain, where the domain comprises n spectrin-like repeats, and contains no more than n spectrin-like repeats, where n is an even number between 4-24, or a nucleic acid encoding a mini-dystrophin peptide. The mini-dystrophin peptide or the polynucleotide encoding it is useful as a medicament, for preparing a drug for therapeutic application and in the preparation of a composition for treatment of muscle disease, e.g. Duchenne's muscular dystrophy (DMD). This sequence represents a human dystrophin polynucleotide CC sequence used in the creation of the mini-dystrophin peptides of the CC invention
 XX SQ Sequence 2691 BP; 860 A; 448 C; 461 G; 922 T; 0 U; 0 Other;
 Query Match 100.0%; Score 137; DB 6; Length 2691;
 Best Local Similarity 100.0%; Pred. No. 3.8e-33;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAATAACGCAATGGCAAGTGGTAGAAGCTGTGAAGCTCAGGTGT 60
 DB 2033 ATTATAAGGAAAAAATAACGCAATGGCAAGTGGTAGAAGCTGTGAAGCTCAGGTGT 1974

QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAAGCTGTGAAGCTCAGGTGT 120
 DB 1973 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAAGCTGTGAAGCTCAGGTGT 1914

QY 121 TTGTGTTTAATTAATT 137
 DB 1913 TTGTGTTTAATTAATT 1897

RESULT 9
 ABK82000/c
 ID ABK82000 standard; DNA; 8689 BP.
 XX AC ABK82000;
 XX DT 13-AUG-2002 (first entry)
 XX DE DNA encoding mini-dystrophin protein deltaH2-R19.
 XX KW Mini-dystrophin peptide; spectrin-like repeat domain; muscle disease;
 XX KW Duchenne's muscular dystrophy; DMD; dystrophin; ds.
 XX OS Homo sapiens.
 XX OS Synthetic.
 XX FN WO200229056-A2.
 XX PD 11-APR-2002.
 XX PF 04-OCT-2001; 2001WO-US0311126.
 XX PR 06-OCT-2000; 2000US-0238848P.
 XX PA (UNMI) UNIV MICHIGAN.
 XX PI Chamberlain JS, Harper SQ;
 XX DR WPI; 2002-435334/46.
 XX CC The invention describes a composition comprising a mini-dystrophin peptide comprising a spectrin-like repeat domain, where the domain comprises n spectrin-like repeats, and contains no more than n spectrin-like repeats, where n is an even number between 4-24, or a nucleic acid encoding a mini-dystrophin peptide. The mini-dystrophin peptide or the polynucleotide encoding it is useful as a medicament, for preparing a drug for therapeutic application and in the preparation of a composition for treatment of muscle disease, e.g. Duchenne's muscular dystrophy (DMD). This sequence represents a mini-dystrophin sequence of the CC invention
 XX SQ Sequence 8689 BP; 2721 A; 1804 C; 1861 G; 2303 T; 0 U; 0 Other;
 Query Match 100.0%; Score 137; DB 6; Length 8689;
 Best Local Similarity 100.0%; Pred. No. 5.2e-33;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAATAACGCAATGGCAAGTGGTAGAAGCTGTGAAGCTCAGGTGT 60
 DB 8031 ATTATAAGGAAAAAATAACGCAATGGCAAGTGGTAGAAGCTGTGAAGCTCAGGTGT 7972

QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAAGCTGTGAAGCTCAGGTGT 120
 DB 7971 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAAGCTGTGAAGCTCAGGTGT 7912

```

QY      121 TTTGATGTTAATTAATT 137
Db      7911 TTGATGTTAATTAATT 7895

RESULT 10
ABK82002/C
ID      ABK82002 standard; DNA; 11443 BP.
XX
XX      AC      ABK82002;
XX      DT      13-AUG-2002 (first entry)
XX
XX      DNA encoding mini-dystrophin protein deltaR9-R16.
XX
XX      Mini-dystrophin peptide; spectrin-like repeat domain; muscle disease;
XX      Duchenne's muscular dystrophy; DMD; dystrophin; ds.
XX
XX      Homo sapiens.
XX      OS
XX      Synthetic.
XX
XX      WO200229056-A2.
XX
XX      11-APR-2002.
XX
XX      04-OCT-2001; 2001WO-US031126.
XX
XX      06-OCT-2000; 2000US-0238848P.
XX
XX      (UNMI ) UNIV MICHIGAN.
XX
XX      Chamberlain JS, Harper SQ;
XX
XX      WPI; 2002-435334/46.
XX
XX      A composition for preparing therapeutic drugs, has a mini-dystrophin
XX      peptide comprising a specific number of spectrin-like repeat domains, or
XX      a nucleic acid sequence encoding the mini-dystrophin peptide.
XX
XX      Disclosure; Fig 17; 145pp; English.
XX
XX
XX      The invention describes a composition comprising a mini-dystrophin
XX      peptide comprising a spectrin-like repeat domain, where the domain
XX      comprises n spectrin-like repeats, and contains no more than n spectrin-
XX      like repeats, where n is an even number between 4-24, or a nucleic acid
XX      encoding a mini-dystrophin peptide. The mini-dystrophin peptide or the
XX      polynucleotide encoding it is useful as a medicament, for preparing a
XX      drug for therapeutic application and in the preparation of a composition
XX      for treatment of muscle disease, e.g. Duchenne's muscular dystrophy
XX      (DMD). This sequence represents a mini-dystrophin sequence of the
XX      invention
XX
XX      Sequence 11443 BP; 3707 A; 2339 C; 2502 G; 2895 T; 0 U; 0 Other;
XX
XX      Query Match      100.0%; Score 137; DB 6; Length 11443;
XX      Best Local Similarity 100.0%; Pred. No. 5.6e-33;
XX      Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY      1 ATTATAAGGAAAGAAAGAAATTAACGCAATGGACAACTGGTGAAGCTGTGAACCTCAGGTGT 60
Db      10785 ATTTAAAGGAAAGAAAGAAATTAACGCAATGGACAACTGGTGAAGCTGTGAACCTCAGGTGT 10726
QY      61 GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGAATAGCATGAGAACCCGTG 120
Db      10725 GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGAATAGCATGAGAACCCGTG 10666
QY      121 TTTGATGTTAATTAATT 137
Db      10665 TTTGATGTTAATTAATT 10649

RESULT 11

```


adrenomedullary	deficiency; severe combined immune deficiency; PAH; beta-chain; haemoglobin gene; beta-thalassaemia; sickle cell disease; low density lipoprotein gene; familial hypercholesterolaemia; hypoxanthine-guanine phosphoribosyltransferase; Lesch-Nyhan syndrome; phenylalanine hydroxylase gene; gene therapy; phenylketonuria; dystrophin gene; muscular dystrophy; cystic fibrosis; immunostimulant; human cystic fibrosis transmembrane conductance regulator gene; antianemic; antilipemic; noctropic; cycostatic; dermatological; human; alpha-1-antitrypsin; lysosomal glucocerebrosidase; ADA; HPRP; lysosomal arylsulphatase A; omithine transcarbamylase; ARSA; OTC; NP; purin nucleoside phosphorylase; gene; ds.
Homo sapiens.	
US2002102731-A1.	
XX	
XX	
XX	
PD	01-AUG-2002.
XX	
PF	12-FEB-2001; 2001US-00782378.
XX	
XX	02-OCT-2000; 2000US-0237747P.
XX	
XX	(UYNY) UNIV NEW YORK STATE RES FOUND.
XX	
XX	Hearing P, Bahou WF, Sandalon Z, Gratenko DV;
XX	WPI; 2002-590619/74.
DR	
XX	
XX	
PT	Producing vector, by introducing vector having nucleotide sequence, adenovirus inverted terminal repeats and packaging sequence, and adenovirus terminal repeat, into cell, and culturing cell.
PT	
XX	Disclosure; Page 122-128; 191pp; English.
XX	
XX	The present invention relates to a new method of producing a vector. The method involves introducing recombinant vector having nucleotide sequence (NS) having 5' and 3' end, left and right inverted terminal repeats of adenovirus flanking NS, adenovirus packaging sequence linked to inverted terminal repeat, and adeno-associated virus terminal repeat linked to 3' end of NS, into cell expressing adenovirus early gene lacking from vector and culturing cell to produce another vector. The method is useful for generating vectors, especially MAD vectors. The method is useful in transferring nucleotide sequences of interest into a cell, for gene transfer applications (e.g. gene therapy) in vitro, ex vivo and in vivo. The nucleotide sequences are useful for treating diseases associated with it, i.e. adenosine deaminase gene associated with adenosine deaminase deficiency with severe combined immune deficiency, beta-chain of haemoglobin gene associated with beta-thalassaemia and sickle cell disease, receptor for low density lipoprotein gene associated with familial hypercholesterolaemia, hypoxanthine-guanine phosphoribosyltransferase associated with Lesch-Nyhan syndrome, phenylalanine hydroxylase (PAH) gene associated with phenylketonuria, dystrophin gene associated with muscular dystrophy, and human cystic fibrosis transmembrane conductance regulator gene associated with cystic fibrosis. The present nucleic acid sequence represents a human disease gene sequence that was used in the methods of the invention
XX	
XX	Sequence 13957 BP; 4602 A; 2781 C; 3122 G; 3452 T; 0 U; 0 Other;
SQ	
	Query Match 100.0%; Score 137; DB 6; Length 13957;
	Best Local Similarity 100.0%; Pred. No. 6e-33;
	Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	1 ATTATTAAGGAAAGAAATTAACCGAATGGCAAGTGGTGAAGCTGTGAACCTAGGTGT 60
DB	13299 ATTATTAAGGAAAGAAATTAACCGAATGGCAAGTGGTGAAGCTGTGAACCTAGGTGT 13240
QY	61 GCACATTTATCAGGACACCCCGAAACCAAGTGGTGAAGTGAAGTGAAGTGAAGCCGTG 120
DB	13239 GCACATTTATCAGGACACCCCGAAACCAAGTGGTGAAGTGAAGTGAAGCCGTG 13180
QY	121 TTGTGATTAATTAAT 137

```

Db      13179 TTTCATGTTAATTAAATT 13163

RESULT 15
ABS70403/c
ID      ABS70403 standard; cDNA; 13977 BP.
XX
XX
AC      ABS70403;
XX
XX      27-NOV-2002 (first entry)
DT
XX
XX      Human bone remodelling gene #60.
DE
XX
XX      Bone remodelling; osteoporosis; human; gene; ss.
KW
XX
XX      Homo sapiens.
OS
XX
XX      US6426186-B1.
PN
XX
XX      30-JUL-2002.
PD
XX
XX      18-JAN-2000; 2000US-00484970.
PP
XX
XX      18-JAN-2000; 2000US-00484970.
PR
XX
XX      (INCY-) INCYTE GENOMICS INC.
PA
XX
XX      Jones KA, Volkmut W, Walker MG;
PI
XX
XX      WPI; 2002-673014/72.
DR
XX
XX      A combination of polynucleotides which are co-expressed with genes known
PT      to be involved in bone remodeling and osteoporosis are useful in an array
PT      for the diagnosis of bone remodeling and osteoporosis associated
PT      disorders.
XX
XX      Claim 1; Col 185-198; 206pp; English.
PS
XX
XX      The invention relates to a combination comprising a number of
CC      substantially purified and isolated polynucleotides which are co-
CC      expressed with genes known to be involved in bone remodeling and
CC      osteoporosis. The invention is used to diagnose disorders associated with
CC      bone remodeling or osteoporosis. ABS70344-ABS70312 represent human bone
CC      remodelling genes of the invention
XX
XX      Sequence 13977 BP; 4596 A; 2765 C; 3120 G; 3453 T; 0 U; 43 Other;
SQ
Query Match      92.0%; Score 126; DB 6; Length 13977;
Best Local Similarity 99.3%; Pred. No. 1.8e-29;
Matches 137; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY      1 ATTATAAGGAAAAGAAAATAATACGCAATGACCAAGTGGTGAAGCTGTGAACCTCAGGTGT 60
13306 ATTATAAGGAAAAGAAAATAATACGCAATGACCAAGTGGTGAAGCTGTGAACCTCAGGTGT 13247
Db
QY      61 GCACAAATTATCAGGACACACCCCAAAACCAAGTCAGGTAGCAATAGCATGAGAG-CCGT 119
13246 GCACAAATTATCAGGACACACCCCAAAACCAAGTCAGGTAGCAATAGCATGAGAGCCCGT 13187
Db
QY      120 GTTCGATGTTAATTAAATT 137
13186 GTTCGATGTTAATTAAATT 13169
Db

Search completed: April 6, 2004, 11:42:37
Job time : 68.812 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 03:22:40 ; Search time 574.36 Seconds
(without alignments)
10338.454 Million cell updates/sec

Title: US-09-966-264d-1
Perfect score: 137
Sequence: 1 attataagaaagaaagaaaa.....gtgttgatgtaattaatt 137

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 3470272 seqs, 21571516995 residues
Total number of hits satisfying chosen parameters: 6940544

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : GenEmbl.*
1: gb.ba.*
2: gb.htg.*
3: gb.in.*
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5: gb.ov.*
6: gb.pat.*
7: gb.ph.*
8: gb.pl.*
9: gb.pr.*
10: gb.ro.*
11: gb.sts.*
12: gb.sy.*
13: gb.un.*
14: gb.vi.*
15: em.ba.*
16: em.fun.*
17: em.hum.*
18: em.in.*
19: em.mu.*
20: em.om.*
21: em.ox.*
22: em.ov.*
23: em.pat.*
24: em.ph.*
25: em.pl.*
26: em.ro.*
27: em.sts.*
28: em.un.*
29: em.vi.*
30: em.htg.hum.*
31: em.htg.inv.*
32: em.htg.other.*
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34: em.htg.pin.*
35: em.htg.tod.*
36: em.htg.nam.*
37: em.htg.vrt.*
38: em.sy.*
39: em.htgo.hum.*
40: em.htgo.mus.*
41: em.htgo.other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
C 1	137	100.0	2148	11	G15848 human STS C
C 2	137	100.0	2563	9	AK129855 Homo sapi
C 3	137	100.0	2691	6	AX538618 Sequence
C 4	137	100.0	4658	9	BC028720 Homo sapi
C 5	137	100.0	8689	6	AX538622 Sequence
C 6	137	100.0	11443	6	AX538624 Sequence
C 7	137	100.0	13957	6	AX409637 Sequence
C 8	137	100.0	13957	6	AX538581 Sequence
C 9	137	100.0	13957	9	M18533 Homo sapien
C 10	137	100.0	98056	9	AC006061 Homo sapi
C 11	126	92.0	13977	6	AR220819 Sequence
C 12	102.8	75.0	13887	4	AF070485 Canis fam
C 13	100.6	73.4	13815	6	AX306153 Sequence
C 14	100.6	73.4	13815	6	AX538582 Sequence
C 15	100.6	73.4	13815	10	MUS2YSA
C 16	100.6	73.4	13907	6	AR093392 Sequence
C 17	100.6	73.4	19307	6	AR142592 Sequence
C 18	100.6	73.4	189131	10	AL645477 Mouse DNA
C 19	100.6	73.4	272578	2	AC108338 Rattus no
C 20	100.6	73.4	279539	2	AC114184 Rattus no
C 21	68.4	49.9	13575	5	GGDYS
C 22	65	47.4	212120	2	AC023414 Homo sapi
C 23	39.8	29.1	172734	10	AL528772 Mouse DNA
C 24	35.8	26.1	94665	3	AC099765 Caenorhab
C 25	35.2	25.7	142000	9	AF005404 Homo sapi
C 26	35.2	25.7	153228	2	AC139129 Mus muscu
C 27	35.2	25.7	153949	9	AC091022 Homo sapi
C 28	35.2	25.7	156863	2	AC069249 Homo sapi
C 29	35.2	25.7	158682	2	AC023083 Homo sapi
C 30	35.2	25.7	183365	2	AC015666 Homo sapi
C 31	35.2	25.7	165789	10	AC111145 Mus muscu
C 32	35.2	25.7	180573	9	AC027689 Homo sapi
C 33	35	25.5	172422	2	BX649289 Danio rer
C 34	35	25.5	231578	2	BX548167 Danio rer
C 35	34.6	25.3	137678	8	AP005183 Oryza sat
C 36	34.6	25.3	138653	8	AP005178 Oryza sat
C 37	34.6	25.3	240279	2	AC126486 Rattus no
C 38	34.6	25.3	270456	2	AC111575 Rattus no
C 39	34.4	25.1	495	8	AY202075 Arabidops
C 40	34.4	25.1	50398	3	AC024771 Caenorhab
C 41	34.4	25.1	106688	8	ATAC011436 Arabidops
C 42	34.4	25.1	206217	2	AC006754 Caenorhab
C 43	34.2	25.0	112944	9	AC107082 Homo sapi
C 44	34.2	25.0	169755	2	AP001199 Homo sapi
C 45	34.2	25.0	171366	9	AC130303 Homo sapi

ALIGNMENTS

RESULT 1
G15848/c
LOCUS human STS CHLC.UTR_01924_M18533.2148 bp DNA linear STS 19-JAN-1996
DEFINITION human STS CHLC.UTR_01924_M18533.2148 bp DNA linear STS 19-JAN-1996
ACCESSION G15848
VERSION G15848.1 GI:1161737
KEYWORDS STS; STS sequence; primer; sequence tagged site.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE
Murray, J., Sheffield, V., Weber, J.L., Duyk, G. and Buetow, K.H.
TITLE Cooperative Human Linkage Center

JOURNAL
COMMENT
Unpublished (1995)
Synonyms: UTR_01924_M18533, CHLC.UTR_01924_M18533.T36152
Contact: Dr. Jeffrey C. Murray
UofI
The University of Iowa
Department of Pediatrics, Iowa City, IA 52242, USA
Tel: (319) 356-3508
Fax: (319) 356-3347
Email: jeff-murray@uiowa.edu
Primer A: AACGCATTTTGGGTTGTTTA
Primer B: GATATCAGCCCAAGGATG
STS size: 189
PCR Profile:

denature: 30 seconds at 94 degrees C
annealing: 75 seconds at 55 degrees C
extension: 15 seconds at 72 degrees C
PCR cycles: 27
extension: 6 minutes at 72 degrees C

Protocol:
Template: 30ng genomic DNA
Primer: each 1.5 pmole
dNTPs: each 200 uM
Taq Polymerase: 0.3 units
Total Vol: 10 ul

Buffer:
MgCl2: 1.5mM
KCl: 50mM
Tris: 10mM
pH: 8.3

Prepared with primer pairs derived from M18533.

FEATURES

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Location/Qualifiers
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
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primer_bind
350..369
primer_bind
complement(519..538)

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Best Local Similarity 100.0%; Pred. No. 7.7e-28;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAATAACGCAATGGACAGTGGTGAAGCTGTGAATCAGGTGT 60
DB 2033 ATTATAAGGAAAAAATAACGCAATGGACAGTGGTGAAGCTGTGAATCAGGTGT 1974
QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAGTGAAGTGAATAATAGCATGAGAACCCGTG 120
DB 1973 GCACAAATTATCAGGAACACCCCAAAACCAAGTGAAGTGAATAATAGCATGAGAACCCGTG 1914
QY 121 TTTCATGTTTAATTAATT 137
DB 1913 TTTCATGTTTAATTAATT 1897

RESULT 2
AK129855/c
LOCUS AK129855 2563 bp mRNA linear PRI 10-SEP-2003
DEFINITION Homo sapiens cDNA FL26345 fis, clone HRT03668.
ACCESSION AK129855
VERSION AK129855.1 GI:34526478
KEYWORDS oligo capping; fis (full insert sequence).
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE
AUTHORS Ota, T., Nakagawa, S., Senoh, A., Mizuguchi, H., Inagaki, H., Suzuki, Y.,
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

Hata, H., Nakagawa, K., Mizuno, S., Morinaga, M., Kawamura, M.,
Sugiyama, T., Irie, R., Otsuki, T., Sato, H., Nishikawa, T.,
Sugiyama, A., Kawakami, B., Nagai, K., Isogai, T. and Sugano, S.
NEDO human cDNA sequencing project
Unpublished
2 (bases 1 to 2563)
Direct Submission
Submitted (31-JUL-2003) Sumio Sugano, Institute of Medical Science,
University of Tokyo, Laboratory of Genome Structure, Human Genome
Center, Shirokane-dai, 4-6-1, Minato-ku, Tokyo 108-8639, Japan
(E-mail: flcdna@ims.u-tokyo.ac.jp, Tel: 81-3-5449-5286,
Fax: 81-3-5449-5416)
NEDO human cDNA sequencing project supported by Ministry of
Economy, Trade and Industry of Japan; cDNA full insert sequencing:
Research Association for Biotechnology (RAB); cDNA library
construction and 5'-end one pass sequencing: Institute of Medical
Science, University of Tokyo, Laboratory of Genome Structure, Human
Genome Center; 3'-end one pass sequencing: RAB; clone selection for
full insert sequencing: RAB and Helix Research Institute.
FEATURES
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/clone="HRT03668"
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/notes="Cloning vector: pME18SFL3"

Query Match 100.0%; Score 137; DB 9; Length 2563;
Best Local Similarity 100.0%; Pred. No. 7.5e-28;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAAAATAACGCAATGGACAGTGGTGAAGCTGTGAATCAGGTGT 60
DB 1905 ATTATAAGGAAAAAATAACGCAATGGACAGTGGTGAAGCTGTGAATCAGGTGT 1846
QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAGTGAAGTGAATAATAGCATGAGAACCCGTG 120
DB 1845 GCACAAATTATCAGGAACACCCCAAAACCAAGTGAAGTGAATAATAGCATGAGAACCCGTG 1786
QY 121 TTTCATGTTTAATTAATT 137
DB 1785 TTTCATGTTTAATTAATT 1769

RESULT 3
AX538618/c
LOCUS AX538618 2691 bp DNA linear PAT 23-NOV-2002
DEFINITION Sequence 38 from Patent WO0229056.
ACCESSION AX538618
VERSION AX538618.1 GI:25271161
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
AUTHORS Chamberlain, J.S. and Harper, S.Q.
TITLE Mini-dystrophin nucleic acid and peptide sequences
JOURNAL Patent: WO 0229056-A 38 11-APR-2002;
THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)
FEATURES
source
1..2691
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"

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Best Local Similarity 100.0%; Pred. No. 7.5e-28;

through the I.M.A.G.E.B. Consortium/LLNL at: <http://image.llnl.gov>
 Series: TRAK Plate: 46 Row: J Column: 3
 This clone was selected for full length sequencing because it
 passed the following selection criteria: matched mRNA gi: 5032296
 This clone has the following problem: The cds is short compared to
 the longest cds in the locus.

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1. .4658
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DXS239, DXS268, DXS269, DXS270, DXS272"
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/db_xref="GI:20379676"
/db_xref="LocusID:1756"

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100.0%; Score 137; DB 9; Length 4658;
1 Similarity 100.0%; Pred. No. 7a-28;
137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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21 TTTGATGTTAATT 137
|||||
64 TTTGATGTTAATT 3848

QY	121	TTTGATGTTAATAATT	137	
Db	13179	TTTGATGTTAATAATT	13163	
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LOCUS		13957 bp	mRNA	linear
DEFINITION				
ACCSSION				
VERSION				
KEYWORDS				
SOURCE				
ORGANISM				
REFERENCE				
AUTHORS				
TITLE				
JOURNAL				
MEDLINE				
PUBMED				
REFERENCE				
AUTHORS				
TITLE				
JOURNAL				
MEDLINE				
PUBMED				
COMMENT				
FEATURES				
source				
gene				
CDS				

121 TTTGATGTTAATAATT 137
13179 TTTGATGTTAATAATT 13163

HUMDYS 13957 bp mRNA linear
Homo sapiens dystrophin (DMD) mRNA, complete cds.
M18533.1 M17154 M18026 M20250
VERSION M18533.1 GI:181856
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 1699)
Koenig,M., Hoffman,E.P., Bertelson,C.J., Monaco,A.P., Feener,C. and Kunkel,L.M.
Complete cloning of the Duchenne muscular dystrophy (DMD) cDNA and preliminary genomic organization of the DMD gene in normal and affected individuals
Cell 50 (3), 509-517 (1987)
87273512
3607877
2 (bases 1678 to 3830)
Hoffman,E.P., Monaco,A.P., Feener,C.C. and Kunkel,L.M.
Conservation of the Duchenne muscular dystrophy gene in mice and humans
Science 238 (4825), 347-350 (1987)
88018015
3659917
3 (bases 1 to 13957)
Koenig,M., Monaco,A.P. and Kunkel,L.M.
The complete sequence of dystrophin predicts a rod-shaped cytoskeletal protein
Cell 53 (2), 219-226 (1988)
88194521
3282674
On May 25, 2000 this sequence version replaced gi:340693.
Draft entry and computer-readable sequence kindly provided by M.Koenig, 01-APR-1988 The severity of muscular dystrophy is determined by the size of the deleted DNA segment. Deletions found in different patients were from positions 302-2200, 473-1168, 1691-1810, and 1169-3011.
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RESULT 10

AC006061
 LOCUS
 DEFINITION Homo sapiens X BAC GSHB-19024 (Genome Systems Human BAC Library)
 AC006061
 ACCESSION complete sequence.
 VERSION
 KEYWORDS
 SOURCE HTG.
 ORGANISM Homo sapiens (human)
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 98056)
 Muzny, D., Arenson, A. D., Bouck, J., Brundage, E., Bunac, C., Chen, Z.,
 Di, W., Ding, Y., Dugan, S., Durbin, J., Forcum, J., Garcia, C.,
 Gorrell, J. H., Gorrell, L. L., Hernandez, J., Jackson, L.,
 Kondarewski, N., Leal, B., Lichtarge, O., Liu, W., Logan, O., Lu, J.,
 Martinez, C., Oswald, G., Pampall, L. R., Parish, B. J., Perez, L.,
 Rashid, N. D., Rives, C., Scherer, S. E., Shen, H., Simon, M., Vo, Q.,
 Williamson, A., Worley, K. C., Yu, W., Zhou, X., Nelson, D. and
 Gibbs, R. A.
 Direct Submission
 Unpublished
 2 (bases 1 to 98056)
 Worley, K. C.
 Direct Submission
 Submitted (26-NOV-1998) Molecular and Human Genetics, Baylor
 College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 3 (bases 1 to 98056)
 Worley, K. C.
 Direct Submission
 Submitted (30-JAN-1999) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 4 (bases 1 to 98056)
 Worley, K. C.
 Direct Submission
 Submitted (02-FEB-1999) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 5 (bases 1 to 98056)
 Worley, K. C.
 Direct Submission
 Submitted (04-FEB-1999) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 6 (bases 1 to 98056)
 Worley, K. C.
 Direct Submission
 Submitted (28-MAR-2001) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 7 (bases 1 to 98056)
 Worley, K. C.
 Direct Submission
 Submitted (07-MAR-2002) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 8 (bases 1 to 98056)
 Worley, K. C.
 Direct Submission
 Submitted (01-MAY-2002) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 INFORMATION: <http://gc.bcm.tmc.edu:8088/home.html> or email
 gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the
 entire insert of this clone. Overlapping regions of clones are only
 sequenced and submitted once, so the sequence for the remainder of
 the insert may be found in the record for the adjacent clones.

Overlapping clones are noted at the beginning and end of the
 Features listing.

ANNOTATION OF FEATURES:

STSS are identified using ePCR (Genome Res. 7:541-550) searches
 of a local database that includes entries from dbSTS, GDB, and
 local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green,
 unpublished) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST
 (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the
 EST and cDNA sequences. Genes demonstrate at least two exons
 flanked by consensus splice sites that maintained sequence
 continuity across the splice junctions. Sequences that are not
 identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum
 standard of double strand coverage with a minimum of 2 clones, and 2
 reads with no ambiguities or 2 chemistries with a minimum of 2
 clones and 3 reads with no ambiguities. If the sequence quality for
 a region does not meet this standard, it will be indicated in the
 annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality
 standards - estimated error rate less than 1 per 10,000 bases.

FEATURES	source	Location/Qualifiers
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	1. 1999	
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	complement(518..570)	
repeat_region		/rpt_family="(GGA)n"
	571..729	
repeat_region		/rpt_family="L1MC4"
	730..987	
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	complement(3626..3656)	
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	5702..7108	
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	complement(9069..9144)	
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	complement(9254..9351)	
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Db 13204 CGATTATAGGAACACCCCAAAACCAAGTGAAGTGAAGTGAATAGCATGAGAACCGCTGTTT 13145
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RESULT 13
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LOCUS AX306153
DEFINITION Sequence 904 from Patent WO0188188.
ACCESSION AX306153
VERSION AX306153.1 GI:17645441
KEYWORDS Mus musculus (house mouse)
SOURCE Mus musculus
ORGANISM Mus musculus
Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1
AUTHORS Ishikawa, K., Asai, S., Takahashi, Y., Nagata, T. and Ishii, Y.
TITLE Method for examining ischemic conditions
JOURNAL Patent: WO 0188188-A 904 22-NOV-2001;
SCHOOL Juridical Person Nihon University (JP)
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source 1. 13815
/organism="Mus musculus"
/mol_type="unassigned DNA"
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Db 13073 GATGTTAAAT 13063

RESULT 14
AX338582/c 13815 bp DNA linear PAT 23-NOV-2002
LOCUS AX338582
DEFINITION Sequence 2 from Patent WO0229056.
ACCESSION AX338582
VERSION AX338582.1 GI:25271088
KEYWORDS Mus musculus (house mouse)
SOURCE Mus musculus
ORGANISM Mus musculus
Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1
AUTHORS Chamberlain, J.S. and Harper, S.Q.
TITLE Mini-dystrophin nucleic acid and peptide sequences
JOURNAL Patent: WO 0229056-A 2 11-APR-2002;
THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)
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Best Local Similarity 85.5%; Pred. No. 8.2e-18;
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QY 124 GATGTTAATTA 134

Db 13073 GATGTTAATTA 13063

RESULT 15

MUSDYSA 13815 bp mRNA linear ROD 24-JUN-1996

LOCUS Mouse dystrophin mRNA, complete cds.

DEFINITION M68859

ACCESSION M68859.1 GI:1388025

KEYWORDS dystrophin.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

REFERENCE 1 (sites)

AUTHORS Koenig,M., Hoffman,E.P., Bertelson,C.J., Monaco,A.P., Feener,C. and Kunkel,L.M.

TITLE Complete cloning of the Duchenne muscular dystrophy (DMD) cDNA and preliminary genomic organization of the DMD gene in normal and affected individuals

JOURNAL Cell 50 (3), 509-517 (1987)

MEDLINE 87273512

PUBMED 3607877

REFERENCE 2 (sites)

AUTHORS Hoffman,E.P., Monaco,A.P., Feener,C.C. and Kunkel,L.M.

TITLE Conservation of the Duchenne muscular dystrophy gene in mice and humans

JOURNAL Science 238 (4825), 347-350 (1987)

MEDLINE 88018015

PUBMED 3659917

REFERENCE 3 (bases 1 to 13815)

AUTHORS Bies,R.D., Phelps,S.F., Cortez,M.D., Roberts,R., Caskey,C.T. and Chamberlain,J.S.

TITLE Human and murine dystrophin mRNA transcripts are differentially expressed during skeletal muscle, heart, and brain development

JOURNAL Nucleic Acids Res. 20 (7), 1725-1731 (1992)

MEDLINE 92253376

PUBMED 1579466

REFERENCE 4 (sites)

AUTHORS Im,W.B., Phelps,S.F., Copen,E.H., Adams,E.G., Slighoton,J.L. and Chamberlain,J.S.

TITLE Differential expression of dystrophin isoforms in multiple strains of mdx mice

JOURNAL Unpublished

REFERENCE 5 (bases 1 to 13815)

AUTHORS Chamberlain,J.S., Pearlman,J.A., Muzny,D.M., Civetello,A., Farwell,N.J., Malek,R., Powaser,P., Reeves,A.A., Lee,C.C. and Caskey,C.T.

TITLE Direct Submission

JOURNAL Submitted (01-JAN-1987)

COMMENT On Jun 24, 1996 this sequence version replaced gi:1387997.

FEATURES

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mRNA

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conflict

CDS

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221. .11257

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 11:33:41 ; Search time 342.077 Seconds
(without alignments)
1500.955 Million cell updates/sec

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Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2470632 seqs, 1873875610 residues

Total number of hits satisfying chosen parameters: 4941264

Minimum DB seq length: 0
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Listing first 45 summaries

Database : Published Applications_NA:*

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6	137	100.0	996	9	US-09-966-264-2
7	137	100.0	2691	15	US-10-149-736-38
8	137	100.0	8689	15	US-10-149-736-42
9	137	100.0	11443	15	US-10-149-736-44
10	137	100.0	13957	9	US-09-782-378A-22
11	137	100.0	13957	9	US-09-880-107-2284
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C 21	31.2	22.8	1977	10	US-09-873-367C-994	Sequence 994, App
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C 23	31.2	22.8	1977	12	US-10-342-887-382	Sequence 382, App
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C 28	30.6	22.3	539	9	US-09-917-800A-153	Sequence 153, App
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C 32	30.4	22.2	704	15	US-10-027-632-17066	Sequence 17066, A
C 33	30.4	22.2	763	15	US-10-027-632-126517	Sequence 126517,
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C 35	30	21.9	4974	9	US-09-906-209-17	Sequence 17, Appli
C 36	30	21.9	6611	10	US-09-934-900-26	Sequence 26, Appli
C 37	30	21.9	7701	15	US-10-459-159-1	Sequence 1, Appli
C 38	30	21.9	7701	15	US-10-427-570A-9	Sequence 9, Appli
C 39	30	21.9	99014	9	US-09-880-107-3428	Sequence 3428, Ap
C 40	29.8	21.8	1015	14	US-10-259-165-589	Sequence 589, App
C 41	29.8	21.8	67191	14	US-10-105-612-1	Sequence 1, Appli
C 42	29.6	21.6	277	12	US-10-424-599-44445	Sequence 44445, A
C 43	29.6	21.6	382	12	US-10-424-599-45074	Sequence 45074, A
C 44	29.4	21.5	575	9	US-09-920-300A-4	Sequence 4, Appli
C 45	29.4	21.5	575	13	US-10-033-528-4	Sequence 4, Appli

ALIGNMENTS

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; APPLICANT: Barber, Elizabeth K
; TITLE OF INVENTION: Gene Expression Control Element DNA
; FILE REFERENCE: 896034605001
; CURRENT APPLICATION NUMBER: US/09/966,264
; CURRENT FILING DATE: 2001-09-28
; PRIOR APPLICATION NUMBER: US 60/237,079
; PRIOR FILING DATE: 2000-09-30
; NUMBER OF SEQ ID NOS: 33
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; SEQ ID NO 1
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; FEATURE:
; NAME/KEY: exon
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; OTHER INFORMATION:
; NAME/KEY: polyA site
; LOCATION: (130)..(135)
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; APPLICANT: McNeill, Patricia Dianne
; APPLICANT: Corixa Corporation
; TITLE OF INVENTION: Compositions and Methods for the Detection, Diagnosis and Therapy
; TITLE OF INVENTION: Hematological Malignancies
; FILE REFERENCE: 014058-014402US
; CURRENT APPLICATION NUMBER: US/10/057,475B
; CURRENT FILING DATE: 2002-01-22
; PRIOR APPLICATION NUMBER: US 60/186,126
; PRIOR FILING DATE: 2000-03-01
; PRIOR APPLICATION NUMBER: US 60/190,479
; PRIOR FILING DATE: 2000-03-17
; PRIOR APPLICATION NUMBER: US 60/200,545
; PRIOR FILING DATE: 2000-04-27
; PRIOR APPLICATION NUMBER: US 60/200,303
; PRIOR FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: US 60/200,779
; PRIOR FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: US 60/200,999
; PRIOR FILING DATE: 2000-05-01
; PRIOR APPLICATION NUMBER: US 60/202,084
; PRIOR FILING DATE: 2000-05-04
; PRIOR APPLICATION NUMBER: US 60/206,201
; PRIOR FILING DATE: 2000-05-22
; PRIOR APPLICATION NUMBER: US 60/218,950
; PRIOR FILING DATE: 2000-07-14
; PRIOR APPLICATION NUMBER: US 60/222,903
; PRIOR FILING DATE: 2000-08-03
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; Remaining Prior Application data removed - See File Wrapper or PALM.
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; SOFTWARE: FastSeq for Windows Version 3.0
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; LENGTH: 430
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; ORGANISM: Homo sapiens
US-10-057-475B-3505

Query Match      100.0%; Score 137; DB 15; Length 430;
Best Local Similarity 100.0%; Pred. No. 3.1e-33;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAATAACGCAATGGACAAGTGGTGAAGCTGTGAAGTCAAGTGT 60
    |||||||
Db 287 ATTATAAGGAAAAAGAAATAACGCAATGGACAAGTGGTGAAGCTGTGAAGTCAAGTGT 228

QY 61 GCACATTATCAGGAACACCCCAACCAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGT 120
    |||||||
Db 227 GCACATTATCAGGAACACCCCAACCAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGT 168

QY 121 TTGTGTTTAATTAATT 137
    |||||||
Db 167 TTGTGTTTAATTAATT 151

RESULT 5
US-10-154-884B-3505/c
; Sequence 3505, Application US/10154884B
; Publication No. US2004000551A1
; GENERAL INFORMATION:
; APPLICANT: Gaiger, Alexander
; APPLICANT: Algate, Paul A.
; APPLICANT: Mannion, Jane
; APPLICANT: Retter, Marc W.
; APPLICANT: Corixa Corporation
; TITLE OF INVENTION: Compositions and Methods for the Detection, Diagnosis and Therapy
; TITLE OF INVENTION: Hematological Malignancies
; FILE REFERENCE: 014058-013521US
; CURRENT APPLICATION NUMBER: US/10/154,884B
; CURRENT FILING DATE: 2002-05-23
; PRIOR APPLICATION NUMBER: US 60/186,126
; PRIOR FILING DATE: 2000-03-01
; PRIOR APPLICATION NUMBER: US 60/190,479
; PRIOR FILING DATE: 2000-03-17
; PRIOR APPLICATION NUMBER: US 60/200,545
```

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; PRIOR FILING DATE: 2000-04-27
; PRIOR APPLICATION NUMBER: US 60/200,303
; PRIOR FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: US 60/200,779
; PRIOR FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: US 60/200,999
; PRIOR FILING DATE: 2000-05-01
; PRIOR APPLICATION NUMBER: US 60/202,084
; PRIOR FILING DATE: 2000-05-04
; PRIOR APPLICATION NUMBER: US 60/206,201
; PRIOR FILING DATE: 2000-05-22
; PRIOR APPLICATION NUMBER: US 60/218,950
; PRIOR FILING DATE: 2000-07-14
; PRIOR APPLICATION NUMBER: US 60/222,903
; PRIOR FILING DATE: 2000-08-03
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 11290
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 3505
; LENGTH: 430
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-154-884B-3505

Query Match      100.0%; Score 137; DB 15; Length 430;
Best Local Similarity 100.0%; Pred. No. 3.1e-33;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAATAACGCAATGGACAAGTGGTGAAGCTGTGAAGTCAAGTGT 60
    |||||||
Db 287 ATTATAAGGAAAAAGAAATAACGCAATGGACAAGTGGTGAAGCTGTGAAGTCAAGTGT 228

QY 61 GCACATTATCAGGAACACCCCAACCAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGT 120
    |||||||
Db 227 GCACATTATCAGGAACACCCCAACCAAGTGAAGTGAAGTGAAGTGAAGTGAAGTGAAGT 168

QY 121 TTGTGTTTAATTAATT 137
    |||||||
Db 167 TTGTGTTTAATTAATT 151

RESULT 6
US-09-966-264-2
; Sequence 2, Application US/09966264
; Patent No. US20020099015A1
; GENERAL INFORMATION:
; APPLICANT: Barber, Elizabeth K
; TITLE OF INVENTION: Gene Expression Control Element DNA
; FILE REFERENCE: 896034605001
; CURRENT APPLICATION NUMBER: US/09/966,264
; CURRENT FILING DATE: 2001-09-28
; PRIOR APPLICATION NUMBER: US 60/237,079
; PRIOR FILING DATE: 2000-09-30
; NUMBER OF SEQ ID NOS: 33
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 996
; TYPE: DNA
; ORGANISM: human
; FEATURE:
; NAME/KEY: exon
; LOCATION: (1)..(996)
; OTHER INFORMATION:
; NAME/KEY: misc feature
; LOCATION: (710)..(996)
; OTHER INFORMATION: Nucleotides 710-996 are homologous to a portion of human dystroph
; OTHER INFORMATION: in DNA in the region of exon 79 except that nucleotides 860-996 a
; OTHER INFORMATION: re inverted in comparison to the orientation of the same sequence
; OTHER INFORMATION: in the dystrophin DNA
US-09-966-264-2

Query Match      100.0%; Score 137; DB 9; Length 996;
Best Local Similarity 100.0%; Pred. No. 4.3e-33;
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Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAGAAATACCAATGACCAAGTGTGAACTGAGCTGAACTCAGGTGT 60
DB 860 ATTATAAGGAAAGAAATACCAATGACCAAGTGTGAACTGAGCTGAACTCAGGTGT 919
QY 61 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGGTAGAAATAGCATGAGAACCCGTG 120
DB 920 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGGTAGAAATAGCATGAGAACCCGTG 979
QY 121 TTGTGATGTTAAATTAATT 137
DB 980 TTGTGATGTTAAATTAATT 996

RESULT 7
US-10-149-736-38/c
; Sequence 38, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; PRIOR FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 38
; LENGTH: 2691
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-149-736-38

Query Match 100.0%; Score 137; DB 15; Length 2691;
Best Local Similarity 100.0%; Pred. No. 6.3e-33;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAGAAATACCAATGACCAAGTGTGAACTGAGCTGAACTCAGGTGT 60
DB 2033 ATTATAAGGAAAGAAATACCAATGACCAAGTGTGAACTGAGCTGAACTCAGGTGT 1974
QY 61 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGGTAGAAATAGCATGAGAACCCGTG 120
DB 1973 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGGTAGAAATAGCATGAGAACCCGTG 1914
QY 121 TTGTGATGTTAAATTAATT 137
DB 1913 TTGTGATGTTAAATTAATT 1897

RESULT 8
US-10-149-736-42/c
; Sequence 42, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; PRIOR FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 42

Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAGAAATACCAATGACCAAGTGTGAACTGAGCTGAACTCAGGTGT 60
DB 10785 ATTATAAGGAAAGAAATACCAATGACCAAGTGTGAACTGAGCTGAACTCAGGTGT 10726
QY 61 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGGTAGAAATAGCATGAGAACCCGTG 120
DB 10725 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGGTAGAAATAGCATGAGAACCCGTG 10666
QY 121 TTGTGATGTTAAATTAATT 137
DB 10665 TTGTGATGTTAAATTAATT 10649

RESULT 10
US-09-782-378A-22/c
; Sequence 22, Application US/09782378A
; Patent No. US20020102731A1
; GENERAL INFORMATION:
; APPLICANT: Hearing, Patrick
; APPLICANT: Bahou, Wadie
```

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; LENGTH: 8689
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-10-149-736-42

Query Match 100.0%; Score 137; DB 15; Length 8689;
Best Local Similarity 100.0%; Pred. No. 9.8e-33;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAGAAATACCAATGACCAAGTGTGAACTGAGCTGAACTCAGGTGT 60
DB 8031 ATTATAAGGAAAGAAATACCAATGACCAAGTGTGAACTGAGCTGAACTCAGGTGT 7972
QY 61 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGGTAGAAATAGCATGAGAACCCGTG 120
DB 7971 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGGTAGAAATAGCATGAGAACCCGTG 7912
QY 121 TTGTGATGTTAAATTAATT 137
DB 7911 TTGTGATGTTAAATTAATT 7895

RESULT 9
US-10-149-736-44/c
; Sequence 44, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; PRIOR FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 44
; LENGTH: 11443
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-10-149-736-44

Query Match 100.0%; Score 137; DB 15; Length 11443;
Best Local Similarity 100.0%; Pred. No. 1.1e-32;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAGAAATACCAATGACCAAGTGTGAACTGAGCTGAACTCAGGTGT 60
DB 10785 ATTATAAGGAAAGAAATACCAATGACCAAGTGTGAACTGAGCTGAACTCAGGTGT 10726
QY 61 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGGTAGAAATAGCATGAGAACCCGTG 120
DB 10725 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGGTAGAAATAGCATGAGAACCCGTG 10666
QY 121 TTGTGATGTTAAATTAATT 137
DB 10665 TTGTGATGTTAAATTAATT 10649

RESULT 10
US-09-782-378A-22/c
; Sequence 22, Application US/09782378A
; Patent No. US20020102731A1
; GENERAL INFORMATION:
; APPLICANT: Hearing, Patrick
; APPLICANT: Bahou, Wadie
```

```

; APPLICANT: Sandalon, Ziv
; APPLICANT: Gnatenko, Dmitri
; TITLE OF INVENTION: Adenoviral Vectors
; FILE REFERENCE: STONY-04970
; CURRENT APPLICATION NUMBER: US/09/782,378A
; PRIOR FILING DATE: 2001-02-12
; PRIOR APPLICATION NUMBER: 60/237,747
; PRIOR FILING DATE: 2000-10-02
; NUMBER OF SEQ ID NOS: 27
; SOFTWARE: Patent in version 3.0
; SEQ ID NO 22
; LENGTH: 13957
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-782-378A-22

Query Match 100.0%; Score 137; DB 9; Length 13957;
Best Local Similarity 100.0%; Pred. No. 1.2e-32;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAAAATACGCAATGGACAAGTGTGAGCTGTGAATCTCAGGTGT 60
DB 13299 ATTATAAGGAAAAAGAAAAATACGCAATGGACAAGTGTGAGCTGTGAATCTCAGGTGT 13240

QY 61 GCACATTTATCAGGAACACCCCAAAACCAAGTGGAGTAGAATAGCATGAGAGCCGTG 120
DB 13239 GCACATTTATCAGGAACACCCCAAAACCAAGTGGAGTAGAATAGCATGAGAGCCGTG 13180

QY 121 TTGTGTTTAATTAATT 137
DB 13179 TTGTGTTTAATTAATT 13163

RESULT 11
US-09-880-107-2284/c
; Sequence 2284, Application US/09880107
; Patent No. US20020142981A1
; GENERAL INFORMATION:
; APPLICANT: Horne, Darci T.
; APPLICANT: Vockley, Joseph G.
; APPLICANT: Scherf, Uwe
; APPLICANT: Gene Logic, Inc.
; TITLE OF INVENTION: Gene Expression Profiles in Liver Cancer
; CURRENT APPLICATION NUMBER: US/09/880,107
; CURRENT FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/211,379
; PRIOR FILING DATE: 2000-06-14
; PRIOR APPLICATION NUMBER: US 60/237,054
; PRIOR FILING DATE: 2000-10-02
; NUMBER OF SEQ ID NOS: 3950
; SOFTWARE: Patent in Ver. 2.1
; SEQ ID NO 2284
; LENGTH: 13957
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: Genbank Accession No. US20020142981A1 M18533
US-09-880-107-2284

Query Match 100.0%; Score 137; DB 9; Length 13957;
Best Local Similarity 100.0%; Pred. No. 1.2e-32;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAAAATACGCAATGGACAAGTGTGAGCTGTGAATCTCAGGTGT 60
DB 13299 ATTATAAGGAAAAAGAAAAATACGCAATGGACAAGTGTGAGCTGTGAATCTCAGGTGT 13240

QY 61 GCACATTTATCAGGAACACCCCAAAACCAAGTGGAGTAGAATAGCATGAGAGCCGTG 120
DB 13239 GCACATTTATCAGGAACACCCCAAAACCAAGTGGAGTAGAATAGCATGAGAGCCGTG 13180

QY 121 TTGTGTTTAATTAATT 137
DB 13179 TTGTGTTTAATTAATT 13163

RESULT 13
US-10-342-887-434/c
; Sequence 434, Application US/10342887
; Publication No. US20040058340A1
; GENERAL INFORMATION:
; APPLICANT: Dai, Hongyue
; APPLICANT: He, Yudong
; APPLICANT: Linsley, Peter S.
; APPLICANT: Mao, Mao
; APPLICANT: Roberts, Christopher J.
; APPLICANT: Van 't Veer, Laura Johanna
; APPLICANT: Van de Vijver, Marc J.
; APPLICANT: Bernards, Rene
; TITLE OF INVENTION: Diagnosis and Prognosis of Breast Cancer Patients
; FILE REFERENCE: 9301-188-999
; CURRENT APPLICATION NUMBER: US/10/342,887
; CURRENT FILING DATE: 2003-01-15
; PRIOR APPLICATION NUMBER: 60/298,918
; PRIOR FILING DATE: 2001-06-18
; PRIOR APPLICATION NUMBER: 60/380,710
; PRIOR FILING DATE: 2002-05-14
; PRIOR APPLICATION NUMBER: 10/172,118
; PRIOR FILING DATE: 2002-06-14
; NUMBER OF SEQ ID NOS: 2699
; SEQ ID NO 434
; LENGTH: 14069
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-342-887-434
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	Query Match	100.0%;	Score 137;	DB 12;	Length 14069;
	Best Local Similarity	100.0%;	Pred. No. 1.2e-32;		
	Matches 137;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
Qy	1	ATTATTAACGGAAAAAGAAATAATACCGAATGGCAAGTGGTGAAGCTGTGAACCTCAGGGTGT	60		
Db	13411	ATTATTAAGGAAAAAGAAATAATACCGAATGGCAAGTGGTGAAGCTGTGAACCTCAGGGTGT	13352		
Qy	61	GCACAAATTATCAGGAACACCCCAAAACCAAGTGAAGGTAGAAATAGCATGAGAAGCCGTG	120		
Db	13351	GCACAAATTATCAGGAACACCCCAAAACCAAGTGAAGGTAGAAATAGCATGAGAAGCCGTG	13292		
Qy	121	TTTGATGTTAAATTAATT	137		
Db	13291	TTTGATGTTAAATTAATT	13275		

RESULT 14
US-10-342-887-981/c
; Sequence 981, Application US/10342887
; Publication No. US20040058340A1
; GENERAL INFORMATION:
; APPLICANT: Dai, Hongyue
; APPLICANT: He, Yudong
; APPLICANT: Linsley, Peter S.
; APPLICANT: Mao, Mao
; APPLICANT: Roberts, Christopher J.
; APPLICANT: Van 't Veer, Laura Johanna
; APPLICANT: Van de Vijver, Marc J.
; APPLICANT: Bernards, Rene
; TITLE OF INVENTION: Diagnosis and Prognosis of Breast Cancer Patients
; FILE REFERENCE: 9301-188-999
; CURRENT APPLICATION NUMBER: US/10/342,887
; CURRENT FILING DATE: 2003-01-15
; PRIOR APPLICATION NUMBER: 60/298,918
; PRIOR FILING DATE: 2001-06-18
; PRIOR APPLICATION NUMBER: 60/380,710
; PRIOR FILING DATE: 2002-05-14
; PRIOR APPLICATION NUMBER: 10/172,118
; PRIOR FILING DATE: 2002-06-14
; NUMBER OF SEQ ID NOS: 2699
; SEQ ID NO 981
; LENGTH: 14082
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-342-887-981

	Query Match	100.0%;	Score 137;	DB 12;	Length 14082;
	Best Local Similarity	100.0%;	Pred. No. 1.2e-32;		
	Matches 137;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
QY	1	ATTATATAGGAAAAAGAAATATACGCAATGCGCAAGTGGTGAAGCTGTGAACCTCAGGTGT	60		
Db	13424	ATTATATAGGAAAAAGAAATATACGCAATGCGCAAGTGGTGAAGCTGTGAACCTCAGGTGT	13365		
QY	61	GCACAAATTATCAGGACACACCCCAAAACCAAAAGTGAGGTAGAAATAGCATGAGAAGCCGTG	120		
Db	13364	GCACAAATTATCAGGACACACCCCAAAACCAAAAGTGAGGTAGAAATAGCATGAGAAGCCGTG	13305		
QY	121	TTTGATGTTTAATTAATT	137		
Db	13304	TTTGATGTTTAATTAATT	13288		

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RESULT 15
US-10-341-434-108/C
; Sequence 108, application US/10341434
; Publication No. US20030215835A1
; GENERAL INFORMATION:
; APPLICANT: Origene Technologies
; TITLE OF INVENTION: Differentially Regulated Prostate Cancer Genes
; FILE REFERENCE: 9U 204 205 R1

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; CURRENT APPLICATION NUMBER: US/10/341,434
; CURRENT FILING DATE: 2003-07-18
; PRIOR APPLICATION NUMBER: US 60/348,164
; PRIOR FILING DATE: 2002-01-15
; PRIOR APPLICATION NUMBER: US 60/348,119
; PRIOR FILING DATE: 2002-01-15
; NUMBER OF SEQ ID NOS: 238
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 108
; LENGTH: 14082
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (703)..(11388)
; OTHER INFORMATION:
; US-10-341-434-108

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	Query Match	Best Local Similarity	100.0%;	Score 137;	DB 15;	Length 14082;
	Matches 137;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;	
Qy	1	ATTATTAAGGAAAGAAAGAAATTAACCCATGACACAAAGTCGCTGAACCTGTGAACCTCAGGTGT	60			
Db	13424	ATTATTAAGGAAAGAAAGAAATTAACCCATGACACAAAGTCGCTGAACCTGTGAACCTCAGGTGT	13365			
Qy	61	GCACAAATTATCAGGACACACCCCAAAACCAAAAGTCAGGTGAGAAATAGCATGAGAAAGCCGTG	120			
Db	13364	GCACAAATTATCAGGACACACCCCAAAACCAAAAGTCAGGTGAGAAATAGCATGAGAAAGCCGTG	13305			
Qy	121	TTTGATGTTAAATTAATT	137			
Db	13304	TTTGATGTTAAATTAATT	13288			

Search completed: April 6, 2004, 15:01:39
Job time : 343.077 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 08:36:01 ; Search time 15.2357 Seconds
(without alignments)
4990.154 Million cell updates/sec

Title: US-09-966-264D-1

Perfect score: 137

Sequence: 1 attataaggaagaaagaaa.....gtgttgatgtaattaatt 137

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Gapop 10.0 , Gapext 1.0

Searched: 682709 seqs, 277475446 residues

Total number of hits satisfying chosen parameters: 1365418

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued Patents NA:*

1: /cgn2_6/ptodata/2/ina/5A COMB.seq:*

2: /cgn2_6/ptodata/2/ina/5B COMB.seq:*

3: /cgn2_6/ptodata/2/ina/6A COMB.seq:*

4: /cgn2_6/ptodata/2/ina/6B COMB.seq:*

5: /cgn2_6/ptodata/2/ina/6C COMB.seq:*

6: /cgn2_6/ptodata/2/ina/backfiles1.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	126	92.0	13977	4	US-09-484-970B-60
C 2	100.6	73.4	19307	3	US-08-836-022A-10
C 3	100.6	73.4	19307	3	US-09-427-048A-10
C 4	28.2	20.6	5326	3	US-09-338-907-124
C 5	28.2	20.6	5326	4	US-09-218-207-124
C 6	28	20.4	38844	4	US-09-734-675-3
C 7	28	20.4	84495	4	US-09-797-906-3
C 8	27.8	20.3	1266	4	US-09-543-681A-2904
C 9	27.8	20.3	2274	3	US-08-423-453-13
C 10	27.8	20.3	2274	3	US-08-423-753-13
C 11	27.8	20.3	2274	3	US-08-716-873-27
C 12	27.8	20.3	2274	3	US-09-368-431-27
C 13	27.8	20.3	2274	4	US-09-414-006-13
C 14	27.8	20.3	2274	4	US-09-447-223-13
C 15	27.8	20.3	2799	2	US-08-874-186-42
C 16	27.8	20.3	7218	1	US-08-232-463-14
C 17	27.6	20.1	1173	4	US-09-107-532A-2915
C 18	27.6	20.1	3705	5	PCT-US96-03940-7
C 19	27.6	20.1	3786	4	US-08-961-527-182
C 20	27.6	20.1	5648	5	PCT-US96-03940-1
C 21	27.6	20.1	18355	4	US-08-956-171E-67
C 22	27.4	20.0	2421	3	US-08-851-843A-51
C 23	27.4	20.0	2421	3	US-08-974-549A-218
C 24	27.4	20.0	2421	3	US-08-854-050-51
C 25	27.4	20.0	2421	4	US-09-430-323-51
C 26	27.4	20.0	2421	4	US-09-402-181B-218
C 27	27.4	20.0	2421	4	US-09-721-456-218

C	28	27.4	20.0	3459	4	US-09-016-434-1363	Sequence 1363, Ap
	29	27.4	20.0	58407	4	US-08-916-421B-2	Sequence 2, Appli
	30	27.2	19.9	770	4	US-09-205-258-161	Sequence 161, App
	31	27.2	19.9	1296	4	US-09-134-001C-1420	Sequence 1420, Ap
	32	27.2	19.9	2508	2	US-08-850-993-1	Sequence 1, Appli
	33	27.2	19.9	13417	2	US-08-637-759B-37	Sequence 37, Appl
	34	27.2	19.9	13417	3	US-08-871-355A-37	Sequence 37, Appl
	35	27.2	19.9	13417	4	US-09-201-945-37	Sequence 37, Appl
	36	26.8	19.6	888	4	US-09-134-001C-1629	Sequence 1629, Ap
	37	26.8	19.6	1321	4	US-09-134-001C-733	Sequence 733, App
	38	26.8	19.6	5873	1	US-07-928-464-4	Sequence 4, Appli
	39	26.8	19.6	5873	5	PCT-US93-07347-4	Sequence 4, Appli
	40	26.8	19.6	5890	1	US-07-928-464-3	Sequence 3, Appli
	41	26.8	19.6	5890	1	US-07-928-464-5	Sequence 5, Appli
	42	26.8	19.6	5890	1	PCT-US93-07347-3	Sequence 3, Appli
	43	26.8	19.6	5890	5	PCT-US93-07347-5	Sequence 5, Appli
	44	26.8	19.6	5890	5	PCT-US93-07347-5	Sequence 5, Appli
	45	26.8	19.6	5890	5	PCT-US93-07347-6	Sequence 6, Appli

ALIGNMENTS

RESULT 1

US-09-484-970B-60/c

; Sequence 60, Application US/09484970B

; Patent No. 6426186

; GENERAL INFORMATION:

; APPLICANT: Jones, Karen A.

; APPLICANT: Volkmut, Wayne

; APPLICANT: Walker, Michael G.

; TITLE OF INVENTION: BONE REMODELING GENES

; FILE REFERENCE: PB-0014 US

; CURRENT APPLICATION NUMBER: US/09/484,970B

; CURRENT FILING DATE: 2000-01-18

; NUMBER OF SEQ ID NOS: 172

; SOFTWARE: PERL Program

; SEQ ID NO 60

; LENGTH: 13977

; TYPE: DNA

; ORGANISM: Homo sapiens

; FEATURE:

; NAME/KEY: misc_feature

; OTHER INFORMATION: Incyte ID No. 6426186 229357.11CB1

; NAME/KEY: unsure

; LOCATION: 11721-11761, 12294, 13969

; OTHER INFORMATION: a, t, c, g, or other

; US-09-484-970B-60

Query Match 92.0%; Score 126; DB 4; Length 13977;
Best Local Similarity 99.3%; Pred. NO. 4.8e-31;
Matches 137; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Qy	1	ATTATAAGGAAAGAAATACGCAATGGCAAGTGGTGAAGCTGCTGAGCTGCACTCAGTGT 60
Db	13306	ATTATAAGGAAAGAAATACGCAATGGCAAGTGGTGAAGCTGCTGAGCTGCACTCAGTGT 13247
Qy	61	GCACATTTATCAGAACACCCCAAAACCAAAATGAGTGAAGTGAATAGCATGAGAAG-CCGT 119
Db	13246	GCACATTTATCAGAACACCCCAAAACCAAAATGAGTGAAGTGAATAGCATGAGAAGCCGT 13187
Qy	120	GTTTGATGTTTAATTAATT 137
Db	13186	GTTTGATGTTTAATTAATT 13169

RESULT 2

US-08-836-022A-10

; Sequence 10, Application US/08836022A

; Patent No. 6001557

; GENERAL INFORMATION:

; APPLICANT: Trustees of the University of Pennsylvania

; APPLICANT: Wilson, James M.

APPLICANT: Fisher, Krishna J.
APPLICANT: Chen, Shu-Jen
TITLE OF INVENTION: Improved Adenovirus Virus and
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Howson and Howson
STREET: Spring House Corporate Cntr, P O Box 457
CITY: Spring House
STATE: Pennsylvania
COUNTRY: USA
ZIP: 19477
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,022A
FILING DATE: 28-OCT-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/331,381
FILING DATE: 28-OCT-1994
ATTORNEY/AGENT INFORMATION:
NAME: Bak, Mary E.
REGISTRATION NUMBER: 31,215
REFERENCE/DOCKET NUMBER: GNVFN.008PCT
TELECOMMUNICATION INFORMATION:
TELEPHONE: 215-540-9200
TELEFAX: 215-540-5818
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 19307 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: unknown
MOLECULE TYPE: CDNA
US-08-836-022A-10
Query Match 73.4%; Score 100.6; DB 3; Length 19307;
Best Local Similarity 85.5%; Pred. No. 8e-23;
Matches 112; Conservative 0; Mismatches 19; Indels 0; Gaps 0;
QY 4 ATAAAGGAAAAAATAACGCAAGTGTGAAGCTGTGAAGCTGCACTCAGGTGTGCA 63
DB 1492 AAAAGGAACTGTGTCCACACACGCAAGTGTGAAGTGTGAAGTGTGAAGTGTGCA 1551
QY 64 CAATTATCAGGAACACCCCAAAACCAAGTGAAGTGAAGTGAAGTGAAGTGTGTTT 123
DB 1552 CAATTACTAGAAACACCCCAAAACCAAGTGAAGTGAAGTGAAGTGAAGTGTGTTT 1611
QY 124 GATGTTAATTA 134
DB 1612 GATGTTAATTA 1622
RESULT 3
US-09-427-048A-10
Sequence 10, Application US/09427048A
Patent No. 6203975
GENERAL INFORMATION:
APPLICANT: Trustees of the University of Pennsylvania
Fisher, Krishna J.
Chen, Shu-Jen
Weitzman, Matthew
TITLE OF INVENTION: Improved Adenovirus Virus and
METHODS OF USE THEREOF
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Howson and Howson
STREET: Spring House Corporate Cntr, P O Box 457

CITY: Spring House
STATE: Pennsylvania
COUNTRY: USA
ZIP: 19477
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/427,048A
FILING DATE: 21-Oct-1999
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,022
FILING DATE: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Bak, Mary E.
REGISTRATION NUMBER: 31,215
REFERENCE/DOCKET NUMBER: GNVFN.008PCT
TELECOMMUNICATION INFORMATION:
TELEPHONE: 215-540-9200
TELEFAX: 215-540-5818
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 19307 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: unknown
MOLECULE TYPE: CDNA
SEQUENCE DESCRIPTION: SEQ ID NO: 10:
US-09-427-048A-10
Query Match 73.4%; Score 100.6; DB 3; Length 19307;
Best Local Similarity 85.5%; Pred. No. 8e-23;
Matches 112; Conservative 0; Mismatches 19; Indels 0; Gaps 0;
QY 4 ATAAAGGAAAAAATAACGCAAGTGTGAAGCTGTGAAGCTGTGAAGTGTGCA 63
DB 1492 AAAAGGAACTGTGTCCACACACGCAAGTGTGAAGTGTGAAGTGTGCA 1551
QY 64 CAATTATCAGGAACACCCCAAAACCAAGTGAAGTGAAGTGAAGTGAAGTGTGTTT 123
DB 1552 CAATTACTAGAAACACCCCAAAACCAAGTGAAGTGAAGTGAAGTGTGTTT 1611
QY 124 GATGTTAATTA 134
DB 1612 GATGTTAATTA 1622
RESULT 4
US-09-338-907-124/c
Sequence 124, Application US/09338907
Patent No. 6265546
GENERAL INFORMATION:
APPLICANT: Cohen, Daniel
APPLICANT: Blumenfeld, Marta
APPLICANT: Ilya, Chumakov
APPLICANT: Bouqueleret, Lydie
TITLE OF INVENTION: PROSTATE CANCER GENE
FILE REFERENCE: GENSET.18CP1CP
CURRENT APPLICATION NUMBER: US/09/338,907
CURRENT FILING DATE: 1999-06-23
EARLIER APPLICATION NUMBER: 08/996,306
EARLIER FILING DATE: 1997-12-22
EARLIER APPLICATION NUMBER: 60/099,658
EARLIER FILING DATE: 1998-09-09
EARLIER APPLICATION NUMBER: 09/218,207
EARLIER FILING DATE: 1998-12-22
NUMBER OF SEQ ID NOS: 578
SOFTWARE: Patent.pm
SEQ ID NO 124
LENGTH: 5324

TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: CDS
LOCATION: 31..588
NAME/KEY: polyA signal
LOCATION: 5297..5302
OTHER INFORMATION: AATAAA
US-09-338-907-124

Query Match 20.6%; Score 28.2; DB 3; Length 5326;
Best Local Similarity 52.8%; Pred. No. 11;
Matches 57; Conservative 1; Mismatches 50; Indels 0; Gaps 0;
QY 11 AAAAGAAATAACGCAATGACCAAGTGTGAAGCTGTGAAGTGTGAAGTGTGACAAATTAT 70
DB 661 ACAGAGAACACGAGCTGGR.GGCGGTGGCAGCTGTATGTCTGTAGTCTTAAGTACT 602
QY 71 CAGGAACACCCCAAAACCAAGTGTAGGTTAGAAATAGCATGAGAACCG 118
DB 601 CAGGAGGACTGCTTTAGCCACGTTGGCGAGCAATGCCTGACTAGCTG 554

RESULT 5

US-09-218-207-124/c
Sequence 124, Application US/09218207
Patent No. 6346381
GENERAL INFORMATION:
APPLICANT: Cohen, Daniel
APPLICANT: Blumenfeld, Marta
APPLICANT: Iliya, Chumakov
APPLICANT: Bougueleret, Lydie
TITLE OF INVENTION: Prostate cancer gene
FILE REFERENCE: GENSET.018CPI
CURRENT APPLICATION NUMBER: US/09/218,207
EARLIER FILING DATE: 1998-12-22
EARLIER APPLICATION NUMBER: 08/996,306
EARLIER FILING DATE: 1997-12-22
EARLIER APPLICATION NUMBER: 60/099,658
EARLIER FILING DATE: 1998-09-09
NUMBER OF SEQ ID NOS: 578
SOFTWARE: Patent.pm
SEQ ID NO 124
LENGTH: 5324
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: CDS
LOCATION: 31..588
NAME/KEY: polyA signal
LOCATION: 5297..5302
OTHER INFORMATION: AATAAA
US-09-218-207-124

Query Match 20.6%; Score 28.2; DB 4; Length 5326;
Best Local Similarity 52.8%; Pred. No. 11;
Matches 57; Conservative 1; Mismatches 50; Indels 0; Gaps 0;
QY 11 AAAAGAAATAACGCAATGACCAAGTGTGAAGCTGTGAAGTGTGACCAATTAT 70
DB 661 ACAGAGAACACGAGCTGGR.GGCGGTGGCAGCTGTATGTCTGTAGTCTTAAGTACT 602
QY 71 CAGGAACACCCCAAAACCAAGTGTAGGTTAGAAATAGCATGAGAACCG 118
DB 601 CAGGAGGACTGCTTTAGCCACGTTGGCGAGCAATGCCTGACTAGCTG 554

RESULT 6

US-09-734-675-3
Sequence 3, Application US/09734675
Patent No. 6365391
GENERAL INFORMATION:
APPLICANT: WEBSTER, Marion et al

TITLE OF INVENTION: ISOLATED HUMAN PROTEASE PROTEINS,
TITLE OF INVENTION: NUCLEIC ACID MOLECULES ENCODING HUMAN PROTEASE PROTEINS, AND
TITLE OF INVENTION: USES THEREOF
FILE REFERENCE: CL000862
CURRENT APPLICATION NUMBER: US/09/734,675
CURRENT FILING DATE: 2000-12-13
NUMBER OF SEQ ID NOS: 4
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 3
LENGTH: 38844
TYPE: DNA
ORGANISM: Human
US-09-734-675-3

Query Match 20.4%; Score 28; DB 4; Length 38844;
Best Local Similarity 50.8%; Pred. No. 23;
Matches 67; Conservative 0; Mismatches 65; Indels 0; Gaps 0;
QY 1 ATTATAAGGMAAAGAAATAACGCAATGGAAGTGTGAAGTGTGAAGTGTGAAGTGTG 60
DB 526 AATATGTATGAGTGAATGAATGATTAATGATTAATGATTAATGATTAATGATTA 585
QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAGTGTAGGTTAGAAATAGCATGAGAACCGGTG 120
DB 586 ACCAAATCAATCGAGAGGACCGAGATAAATTTGTGCTCTAGAGTAAGAGACCTGAG 645
QY 121 TTTGATGTTAAT 132
DB 646 TTTGAGATAACT 657

RESULT 7

US-09-797-906-3/c
Sequence 3, Application US/09797906
Patent No. 6329188
GENERAL INFORMATION:
APPLICANT: Zianghe YAN, Karen A. KETCHUM, Valentina DIFRANCESCO, Ellen M. BEASLEY
TITLE OF INVENTION: ISOLATED HUMAN PROTEASE PROTEINS,
TITLE OF INVENTION: NUCLEIC ACID MOLECULES ENCODING HUMAN PROTEASE PROTEINS, AND
TITLE OF INVENTION: USES THEREOF
FILE REFERENCE: CL001151CIP
CURRENT APPLICATION NUMBER: US/09/797,906
CURRENT FILING DATE: 2001-03-05
NUMBER OF SEQ ID NOS: 5
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 3
LENGTH: 84495
TYPE: DNA
ORGANISM: Human
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)...(84495)
OTHER INFORMATION: n = A,T,C or G
US-09-797-906-3

Query Match 20.4%; Score 28; DB 4; Length 84495;
Best Local Similarity 55.0%; Pred. No. 29;
Matches 55; Conservative 0; Mismatches 45; Indels 0; Gaps 0;
QY 2 TTATAAGGMAAAGAAATAACGCAATGGAAGTGTGAAGTGTGAAGTGTGAACTGAGTGTG 61
DB 9226 TTATAAGGMAAAGAAATAACGCAATGGAAGTGTGAAGTGTGAAGTGTGAACTGAGTGTG 9167
QY 62 CACAATTATCAGGAACACCCCAAAACCAAGTGTAGGTTAGAAATAGCATGAGTAGA 101
DB 9166 CAATAAAGGMAAAGAAATAACGCAATGGAAGTGTGAAGTGTGAAGTGTGAACTGAGTGTG 9127

RESULT 8

US-09-543-681A-2904/c
Sequence 2904, Application US/09543681A
Patent No. 6605709
GENERAL INFORMATION:


```

; APPLICANT: GARY BRETON
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PROTEUS MIRABILIS
; FILE REFERENCE: 2709.1002-001
; CURRENT APPLICATION NUMBER: US/09/543,681A
; PRIOR FILING DATE: 2000-04-05
; PRIOR FILING DATE: 1999-04-09
; NUMBER OF SEQ ID NOS: 8344
; SEQ ID NO 2904
; LENGTH: 1266
; TYPE: DNA
; ORGANISM: Proteus mirabilis
US-09-543-681A-2904

Query Match 20.3%; Score 27.8; DB 4; Length 1266;
Best Local Similarity 55.1%; Pred. No. 9;
Matches 41; Conservative 0; Mismatches 22; Indels 0; Gaps 0;

QY 65 AATTATCAGGACACCCCAAAAGTGGTGAATAGCATGAGAACCGGTGTTG 124
Db 886 ATTATCAGAGAACCCAGCTAAATAAGAGTGGCCAGGATAGCAACCAAGGCATGAGTG 827

QY 125 ATG 127
Db 826 ATG 824

RESULT 9
US-08-492-459-13/c
; Sequence 13, Application US/08492459
; Patent No. 6015689
; GENERAL INFORMATION:
; APPLICANT: Takashi OKADO et al.
; TITLE OF INVENTION: REGULATION OF AUROBASIDIN SENSITIVITY IN FUNGUS
; NUMBER OF SEQUENCES: 36
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Wenderoth, Lind & Ponack
; STREET: 805 Fifteenth Street, N.W., #700
; CITY: Washington
; STATE: D.C.
; COUNTRY: U.S.A.
; ZIP: 20005
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 1.4 mb
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: Wordperfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/492,459
; FILING DATE: June 20, 1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; FILING DATE: May 16, 1994
; APPLICATION NUMBER: 08/243,403
; NAME: Warren M. Cheek, Jr.
; REGISTRATION NUMBER: 33,367
; REFERENCE/DOCKET NUMBER:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-8850
; TELEFAX:
; TELEX:
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2274
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: genomic DNA
US-08-492-459-13

Query Match 20.3%; Score 27.8; DB 3; Length 2274;
Best Local Similarity 55.1%; Pred. No. 9;
Matches 41; Conservative 0; Mismatches 22; Indels 0; Gaps 0;

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Best Local Similarity 57.5%; Pred. No. 11;
Matches 50; Conservative 0; Mismatches 37; Indels 0; Gaps 0;

QY 50 AACTCAGGTGTGCACAATTATCAGGACACCCCAAAAGTGGTGAATAGCAT 109
Db 1848 AACACATTATGTAATAAAGAAAGACACACACACACACACACACACACTTAAATAAAGTGT 1789

QY 110 GAGAAGCGGTGTTGATGTTAATAAT 136
Db 1788 GCCATTCTATATTGATGTTAATAAT 1762

RESULT 10
US-08-423-752-13/c
; Sequence 13, Application US/08423752
; Patent No. 6022349
; GENERAL INFORMATION:
; APPLICANT: Takashi OKADO et al.
; TITLE OF INVENTION: A GENE CODING FOR A PROTEIN REGULATING
; TITLE OF INVENTION: AUROBASIDIN SENSITIVITY
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Wenderoth, Lind & Ponack
; STREET: 805 Fifteenth Street, N.W., #700
; CITY: Washington
; STATE: D.C.
; COUNTRY: U.S.A.
; ZIP: 20005
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 5.25 inch, 500 kb
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: Wordperfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/423,752
; FILING DATE: April 18, 1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/243,403
; FILING DATE: May 16, 1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Warren M. Cheek, Jr.
; REGISTRATION NUMBER: 33,367
; REFERENCE/DOCKET NUMBER:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-8850
; TELEFAX:
; TELEX:
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2274
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: genomic DNA
US-08-423-752-13

Query Match 20.3%; Score 27.8; DB 3; Length 2274;
Best Local Similarity 57.5%; Pred. No. 11;
Matches 50; Conservative 0; Mismatches 37; Indels 0; Gaps 0;

QY 50 AACTCAGGTGTGCACAATTATCAGGACACCCCAAAAGTGGTGAATAGCAT 109
Db 1848 AACACATTATGTAATAAAGAAAGACACACACACACACACACACACTTAAATAAAGTGT 1789

QY 110 GAGAAGCGGTGTTGATGTTAATAAT 136
Db 1788 GCCATTCTATATTGATGTTAATAAT 1762

RESULT 11
US-08-716-873-27/c
; Sequence 27, Application US/08716873

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COUNTRY: U.S.A.
ZIP: 20006
COMPUTER TYPE: FORM:
MEDIUM TYPE: Diskette, 3.5 inch,
COMPUTER: IBM Compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: Wordperfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/414,006
FILING DATE: October 7, 1999
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/492,459
FILING DATE: June 20, 1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/243,403
FILING DATE: May 16, 1994
ATTORNEY/AGENT INFORMATION:
NAME: Warren M. Cheek, Jr.
REGISTRATION NUMBER: 33,367

REFERENCE/DOCKET NUMBER:
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-721-8200
TELEFAX: 202-721-8250
TELEX:
INFORMATION FOR SEQ ID NO: 13:
SEQUENCE CHARACTERISTICS:
LENGTH: 2274
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: genomic DNA
US-09-414-006-13

Query Match 20.3%; Score 27.8; DB 4; Length 2274;
Best Local Similarity 57.5%; Pred. No. 11;
Matches 50; Conservative 0; Mismatches 37; Indels 0; Gaps 0;
QY 50 AACTCAGGTGTCACATTTATCAGGACACCCCAACCAAGTGGTAGAATAGCAT 109
DB 1848 AACACATTAATGTAAAAAAGAAAGACACACACAAAAAAGTAAAAAACTGT 1789
QY 110 GAGAACGCGTTTGTGTTTAATTAAT 136
DB 1788 GCCCATCTATATTGATGGTAATTAAT 1762

RESULT 14
US-09-447-223-13/c
Sequence 13, Application US/09447223
Patent No. 6432664
GENERAL INFORMATION:
APPLICANT: Takashi OKADO et al.
TITLE OF INVENTION: A GENE CODING FOR A PROTEIN REGULATING
AUROBASITIN SENSITIVITY

NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: Wenderoth, Lind & Ponack
STREET: 805 Fifteenth Street, N.W., #700
CITY: Washington
STATE: D.C.
COUNTRY: U.S.A.
ZIP: 20005

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 5.25 inch, 500 kb
COMPUTER: IBM compatible
OPERATING SYSTEM: MS-DOS
SOFTWARE: Wordperfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/447,223
FILING DATE: 23-No. 6432664-1999
CLASSIFICATION: <Unknown>

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/243,403
FILING DATE: May 16, 1994
ATTORNEY/AGENT INFORMATION:
NAME: Warren M. Cheek, Jr.
REGISTRATION NUMBER: 33,367
REFERENCE/DOCKET NUMBER: <Unknown>
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-8850
TELEFAX: <Unknown>
TELEX: <Unknown>

INFORMATION FOR SEQ ID NO: 13:
SEQUENCE CHARACTERISTICS:
LENGTH: 2274
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: genomic DNA
SEQUENCE DESCRIPTION: SEQ ID NO: 13:
US-09-447-223-13

Query Match 20.3%; Score 27.8; DB 4; Length 2274;
Best Local Similarity 57.5%; Pred. No. 11;
Matches 50; Conservative 0; Mismatches 37; Indels 0; Gaps 0;
QY 50 AACTCAGGTGTCACATTTATCAGGACACCCCAACCAAGTGGTAGAATAGCAT 109
DB 1848 AACACATTAATGTAAAAAAGAAAGACACACAAAAAAGTAAAAAACTGT 1789
QY 110 GAGAACGCGTTTGTGTTTAATTAAT 136
DB 1788 GCCCATCTATATTGATGGTAATTAAT 1762

RESULT 15
US-08-874-186-42
Sequence 42, Application US/08874186
Patent No. 5989885
GENERAL INFORMATION:
APPLICANT: Teng, David H-F.
APPLICANT: Tavtigian, Sean V.
APPLICANT: Perry III, William L.
APPLICANT: Skolnick, Mark H.
TITLE OF INVENTION: SPECIFIC MUTATIONS OF MAP KINASE KINASE
IN HUMAN TUMOR CELL LINES IDENTIFY IT AS A TUMOR
SUPPRESSOR IN VARIOUS TYPES OF CANCER
TITLE OF INVENTION: 4 (MKK4)
NUMBER OF SEQUENCES: 96
CORRESPONDENCE ADDRESS:
ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP
STREET: 1201 New York Avenue, N.W., Suite 1000
CITY: Washington
STATE: DC
COUNTRY: U.S.A.
ZIP: 20005

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/874,186
FILING DATE:
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/782,482
FILING DATE: 10-JAN-1997
ATTORNEY/AGENT INFORMATION:
NAME: Saxe, Stephen A.
REGISTRATION NUMBER: 38,609
REFERENCE/DOCKET NUMBER: 24884-121392-01
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-962-4848
TELEFAX: 202-962-8300

INFORMATION FOR SEQ ID NO: 42:
SEQUENCE CHARACTERISTICS:
LENGTH: 2799 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
FEATURE:
NAME/KEY: intron
LOCATION: 1..764
FEATURE:
NAME/KEY: exon
LOCATION: 765..892
FEATURE:
NAME/KEY: intron
LOCATION: 893..2799
US-08-874-186-42

Query Match 20.3%; Score 27.8; DB 2; Length 2799;

Best Local Similarity 57.5%; Pred. No. 12;
Matches 50; Conservative 0; Mismatches 37; Indels 0; Gaps 0;
Qy 18 AAATACCGCAATGGACAACTGGTGAAGCTGTGAACCTCAGGTGTGCACAAATTATCAGGAAC 77
Db 2082 AAATAAATAAATAATTAGACATGGAGTATTTGGAAATGTGATTACATATTTTATTAGGAT 2141
Qy 78 ACCCAAAACCAAGTGAGGTAGAAAT 104
Db 2142 ACCCGAGAATGAAGTGAATTTAT 2168

Search completed: April 6, 2004, 14:14:17
Job time : 16.2357 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 15:01:53 ; Search time 61.7891 Seconds
(without alignments)
8309.593 Million cell updates/sec

Title: US-09-966-264D-1

Perfect score: 137
Sequence: 1 attataaggagaaagaaaa.....gtgttgatgtaattaatt 137

Scoring table: OLIGO NUC

Gapop 60.0 , Gapext 60.0

Searched: 2470632 seqs, 1873875610 residues

Word size : 0

Total number of hits satisfying chosen parameters: 4941264

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : Published Applications NA:*

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- 3: /cgn2_6/ptodata/1/pubpna/US05_NEW_PUB.seq:*
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- 7: /cgn2_6/ptodata/1/pubpna/US08_NEW_PUB.seq:*
- 8: /cgn2_6/ptodata/1/pubpna/US09_PUBCOMB.seq:*
- 9: /cgn2_6/ptodata/1/pubpna/US09A_PUBCOMB.seq:*
- 10: /cgn2_6/ptodata/1/pubpna/US09B_PUBCOMB.seq:*
- 11: /cgn2_6/ptodata/1/pubpna/US09C_PUBCOMB.seq:*
- 12: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq:*
- 13: /cgn2_6/ptodata/1/pubpna/US10A_PUBCOMB.seq:*
- 14: /cgn2_6/ptodata/1/pubpna/US10B_PUBCOMB.seq:*
- 15: /cgn2_6/ptodata/1/pubpna/US10C_PUBCOMB.seq:*
- 16: /cgn2_6/ptodata/1/pubpna/US10_NEW_PUB.seq:*
- 17: /cgn2_6/ptodata/1/pubpna/US60_NEW_PUB.seq:*
- 18: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	137	100.0	137	9	US-09-966-264-1
C 2	137	100.0	430	9	Sequence 1, Appli
C 3	137	100.0	430	14	Sequence 3505, Ap
C 4	137	100.0	430	15	Sequence 3505, Ap
C 5	137	100.0	430	15	Sequence 3505, Ap
C 6	137	100.0	996	9	Sequence 2, Appli
C 7	137	100.0	2691	15	Sequence 38, Appl
C 8	137	100.0	8689	15	Sequence 42, Appl
C 9	137	100.0	11443	15	Sequence 44, Appl
C 10	137	100.0	13957	9	Sequence 22, Appl
C 11	137	100.0	13957	9	Sequence 2284, Ap
C 12	137	100.0	13957	15	Sequence 1, Appli
C 13	137	100.0	14069	12	Sequence 434, App
C 14	137	100.0	14082	12	Sequence 981, App
C 15	137	100.0	14082	15	Sequence 108, App

C 16	13815	15	US-10-149-736-2	Sequence 2, Appli
C 17	22	22	US-09-966-264-13	Sequence 13, Appli
C 18	20	16.1	US-10-131-827-8211	Sequence 8211, Ap
C 19	19	13.9	US-10-027-632-293205	Sequence 293205,
C 20	19	13.9	US-10-027-632-81394	Sequence 81394, A
C 21	19	13.9	US-10-027-632-114-6882	Sequence 6882, Ap
C 22	19	13.9	US-09-373-658-23	Sequence 23, Appli
C 23	19	13.9	US-09-989-687-23	Sequence 23, Appli
C 24	19	13.9	US-10-235-192A-27	Sequence 27, Appli
C 25	18	13.1	US-09-966-264-22	Sequence 22, Appli
C 26	18	13.1	US-10-027-632-274023	Sequence 274023,
C 27	18	13.1	US-10-027-632-280438	Sequence 280438,
C 28	18	13.1	US-10-027-632-280438	Sequence 21513, A
C 29	18	13.1	US-10-027-632-148232	Sequence 148232,
C 30	18	13.1	US-10-027-632-154166	Sequence 154166,
C 31	18	13.1	US-10-027-632-154167	Sequence 154167,
C 32	18	13.1	US-10-027-632-154168	Sequence 154168,
C 33	18	13.1	US-10-027-632-154169	Sequence 154169,
C 34	18	13.1	US-09-770-445-538	Sequence 538, App
C 35	18	13.1	US-10-027-632-258397	Sequence 258397,
C 36	18	13.1	US-10-027-632-258398	Sequence 258398,
C 37	18	13.1	US-10-027-632-102777	Sequence 102777,
C 38	18	13.1	US-09-814-353-19075	Sequence 19075, A
C 39	18	13.1	US-10-135-322-34	Sequence 34, Appli
C 40	18	13.1	US-10-135-322-33	Sequence 33, Appli
C 41	18	13.1	US-10-135-322-13	Sequence 13, Appli
C 42	17	12.4	US-10-085-783A-12712	Sequence 12712, A
C 43	17	12.4	US-10-242-535A-12712	Sequence 12712, A
C 44	17	12.4	US-10-424-599-396	Sequence 396, App
C 45	17	12.4	US-09-918-995-13211	Sequence 13211, A

ALIGNMENTS

RESULT 1
US-09-966-264-1
; Sequence 1, Application US/09966264
; Patent No. US2002009015A1
; GENERAL INFORMATION:
; APPLICANT: Barber, Elizabeth K
; TITLE OF INVENTION: Gene Expression Control Element DNA
; FILE REFERENCE: 8960346US001
; CURRENT APPLICATION NUMBER: US/09/966,264
; CURRENT FILING DATE: 2001-09-28
; PRIOR APPLICATION NUMBER: US 60/237,079
; PRIOR FILING DATE: 2000-09-30
; NUMBER OF SEQ ID NOS: 33
; SOFTWARE: Patent in version 3.1
; SEQ ID NO 1
; LENGTH: 137
; TYPE: DNA
; ORGANISM: human
; FEATURE:
; NAME/KEY: exon
; LOCATION: (3)...(137)
; OTHER INFORMATION:
; NAME/KEY: polyA site
; LOCATION: (130)...(135)
; OTHER INFORMATION:
US-09-966-264-1

Query Match	100.0%	Score 137	DB 9	Length 137
Best Local Similarity	100.0%	Pred. No. 9.5e-61	Mismatches 0	Indels 0
Matches 137	Conservative	0		
QY	1	ATTATAAGGAAAAAAGAAAAATACGCAATGCAAGTGTGAAGCTGTGAACCTCAGGTGT	60	
Db	1	ATTATAAGGAAAAAAGAAAAATACGCAATGCAAGTGTGAAGCTGTGAACCTCAGGTGT	60	
QY	61	GCACAATTATCAGGAACACCCCAACCAAGTAGAGTAAATAGCATGAGAGCCGTC	120	
Db	61	GCACAATTATCAGGAACACCCCAACCAAGTAGAGTAAATAGCATGAGAGCCGTC	120	


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; APPLICANT: McNeill, Patricia Dianne
; APPLICANT: Corixa Corporation
; TITLE OF INVENTION: Compositions and Methods for the Detection, Diagnosis and Therapy
; FILE OF INVENTION: Hematological Malignancies
; FILE REFERENCE: 014058-014402US
; CURRENT APPLICATION NUMBER: US/10/057,475B
; CURRENT FILING DATE: 2002-01-22
; PRIOR APPLICATION NUMBER: US 60/186,126
; PRIOR FILING DATE: 2000-03-01
; PRIOR APPLICATION NUMBER: US 60/190,479
; PRIOR FILING DATE: 2000-03-17
; PRIOR APPLICATION NUMBER: US 60/200,545
; PRIOR FILING DATE: 2000-04-27
; PRIOR APPLICATION NUMBER: US 60/200,303
; PRIOR FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: US 60/200,779
; PRIOR FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: US 60/200,999
; PRIOR FILING DATE: 2000-05-01
; PRIOR APPLICATION NUMBER: US 60/202,084
; PRIOR FILING DATE: 2000-05-04
; PRIOR APPLICATION NUMBER: US 60/206,201
; PRIOR FILING DATE: 2000-05-22
; PRIOR APPLICATION NUMBER: US 60/218,950
; PRIOR FILING DATE: 2000-07-14
; PRIOR APPLICATION NUMBER: US 60/222,903
; PRIOR FILING DATE: 2000-08-03
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 10979
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 3505
; LENGTH: 430
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-057-475B-3505

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Best Local Similarity 100.0%; Pred. No. 1e-60;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAATAACCGAATGCAAGTGGTGAAGCTGTGAAGTGTGAAGTGTGAAGTGT 60
Db 287 ATTATAAGGAAAAAGAAATAACCGAATGCAAGTGGTGAAGTGTGAAGTGTGAAGTGT 228

QY 61 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGTGAAGTGTGAAGTGTGAAGTGT 120
Db 227 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGTGAAGTGTGAAGTGTGAAGTGT 168

QY 121 TTGTGATGTTAATTAATT 137
Db 167 TTGTGATGTTAATTAATT 151

RESULT 5
US-10-154-884B-3505/c
; Sequence 3505, Application US/10154884B
; Publication No. US20040005561A1
; GENERAL INFORMATION:
; APPLICANT: Gaiger, Alexander
; APPLICANT: Algate, Paul A.
; APPLICANT: Mannion, Jane
; APPLICANT: Retter, Marc W.
; APPLICANT: Corixa Corporation
; TITLE OF INVENTION: Compositions and Methods for the Detection, Diagnosis and Therapy
; FILE OF INVENTION: Hematological Malignancies
; FILE REFERENCE: 014058-013521US
; CURRENT APPLICATION NUMBER: US/10/154,884B
; CURRENT FILING DATE: 2002-05-23
; PRIOR APPLICATION NUMBER: US 60/186,126
; PRIOR FILING DATE: 2000-03-01
; PRIOR APPLICATION NUMBER: US 60/190,479
; PRIOR FILING DATE: 2000-03-17
; PRIOR APPLICATION NUMBER: US 60/200,545

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; PRIOR FILING DATE: 2000-04-27
; PRIOR APPLICATION NUMBER: US 60/200,303
; PRIOR FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: US 60/200,779
; PRIOR FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: US 60/200,999
; PRIOR FILING DATE: 2000-05-01
; PRIOR APPLICATION NUMBER: US 60/202,084
; PRIOR FILING DATE: 2000-05-04
; PRIOR APPLICATION NUMBER: US 60/206,201
; PRIOR FILING DATE: 2000-05-22
; PRIOR APPLICATION NUMBER: US 60/218,950
; PRIOR FILING DATE: 2000-07-14
; PRIOR APPLICATION NUMBER: US 60/222,903
; PRIOR FILING DATE: 2000-08-03
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 11290
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 3505
; LENGTH: 430
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-154-884B-3505

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Best Local Similarity 100.0%; Pred. No. 1e-60;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAATAACCGAATGCAAGTGGTGAAGTGTGAAGTGTGAAGTGT 60
Db 287 ATTATAAGGAAAAAGAAATAACCGAATGCAAGTGGTGAAGTGTGAAGTGTGAAGTGT 228

QY 61 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGTGAAGTGTGAAGTGTGAAGTGT 120
Db 227 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGTGAAGTGTGAAGTGTGAAGTGT 168

QY 121 TTGTGATGTTAATTAATT 137
Db 167 TTGTGATGTTAATTAATT 151

RESULT 6
US-09-966-264-2
; Sequence 2, Application US/09966264
; Patent No. US20020099015A1
; GENERAL INFORMATION:
; APPLICANT: Barber, Elizabeth K
; TITLE OF INVENTION: Gene Expression Control Element DNA
; FILE REFERENCE: 996034605001
; CURRENT APPLICATION NUMBER: US/09/966,264
; CURRENT FILING DATE: 2001-09-28
; PRIOR APPLICATION NUMBER: US 60/237,079
; PRIOR FILING DATE: 2000-09-30
; NUMBER OF SEQ ID NOS: 33
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 996
; TYPE: DNA
; ORGANISM: human
; FEATURE:
; NAME/KEY: exon
; LOCATION: (1) ..(996)
; OTHER INFORMATION:
; NAME/KEY: misc feature
; LOCATION: (710) ..(996)
; OTHER INFORMATION: Nucleotides 710-996 are homologous to a portion of human dystroph
; OTHER INFORMATION: in DNA in the region of exon 79 except that nucleotides 860-996 a
; OTHER INFORMATION: re inverted in comparison to the orientation of the same sequence
; OTHER INFORMATION: in the dystrophin DNA
US-09-966-264-2

Query Match      100.0%; Score 137; DB 9; Length 996;
Best Local Similarity 100.0%; Pred. No. 1e-60;

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; APPLICANT: Sandalon, Ziv
; APPLICANT: Gnatenko, Dmitri
; TITLE OF INVENTION: Adenoviral Vectors
; FILE REFERENCE: STONYB-04970
; CURRENT APPLICATION NUMBER: US/09/782,378A
; CURRENT FILING DATE: 2001-02-12
; PRIOR APPLICATION NUMBER: 60/237,747
; PRIOR FILING DATE: 2000-10-02
; NUMBER OF SEQ ID NOS: 27
; SOFTWARE: Patent in version 3.0
; SEQ ID NO 22
; LENGTH: 13957
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-782-378A-22

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Best Local Similarity 100.0%; Pred. No. 1.2e-60;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY      61 GCACAAATTATCAGGACACCCCAAAACCAAAAGTGAGGTAGAAATAGCATGAGAAGCCGTG 120
        |||
Db      13239 GCACAAATTATCAGGACACCCCAAAACCAAAAGTGAGGTAGAAATAGCATGAGAAGCCGTG 13180

QY      121 TTTCATGTTAAATTAATT 137
        |||
Db      13179 TTTCATGTTAAATTAATT 13163

RESULT 11
US-09-880-107-2284/c
; Sequence 2284, Application US/09880107
; Patent No. US20020142981A1
; GENERAL INFORMATION:
; APPLICANT: Horne, Darci T.
; APPLICANT: Vockley, Joseph G.
; APPLICANT: Scherf, Uwe
; APPLICANT: Gene Logic, Inc.
; TITLE OF INVENTION: Gene Expression Profiles in Liver Cancer
; FILE REFERENCE: 44921-5028-WO
; CURRENT APPLICATION NUMBER: US/09/880,107
; CURRENT FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/211,379
; PRIOR FILING DATE: 2000-06-14
; PRIOR APPLICATION NUMBER: US 60/237,054
; PRIOR FILING DATE: 2000-10-02
; NUMBER OF SEQ ID NOS: 3950
; SOFTWARE: Patent in Ver. 2.1
; SEQ ID NO 2284
; LENGTH: 13957
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: Genbank Accession No. US20020142981A1 M18533
US-09-880-107-2284

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Best Local Similarity 100.0%; Pred. No. 1.2e-60;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY      61 GCACAAATTATCAGGACACCCCAAAACCAAAAGTGAGGTAGAAATAGCATGAGAAGCCGTG 120
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Db      13239 GCACAAATTATCAGGACACCCCAAAACCAAAAGTGAGGTAGAAATAGCATGAGAAGCCGTG 13180

QY      121 TTTCATGTTAAATTAATT 137
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Query Match 100.0%; Score 137; DB 12; Length 14069;
Best Local Similarity 100.0%; Pred. No. 1.2e-60;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAAAATACCAATGGACAAGTGTGAAGCTGTGAAGTGTGAAGTGTGT 60
Db 13411 ATTATAAGGAAAAAGAAAAATACCAATGGACAAGTGTGAAGCTGTGAAGTGTGT 13352

QY 61 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGTGAAGTGTGAAGTGTGAAGTGT 120
Db 13351 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGTGAAGTGTGAAGTGTGAAGTGT 13292

QY 121 TTTGATGTTAATTAATT 137
Db 13291 TTTGATGTTAATTAATT 13275

RESULT 14
US-10-342-887-981/c
; Publication 981, Application US/10342887
; Sequence 981, Application US/10342887
; Publication No. US20040058340A1
; GENERAL INFORMATION:
; APPLICANT: Dai, Hongyue
; APPLICANT: He, Yudong
; APPLICANT: Linsley, Peter S.
; APPLICANT: Mao, Mao
; APPLICANT: Roberts, Christopher J.
; APPLICANT: Van 't Veer, Laura Johanna
; APPLICANT: Van de Vijver, Marc J.
; APPLICANT: Bernards, Rene
; TITLE OF INVENTION: Diagnosis and Prognosis of Breast Cancer Patients
; FILE REFERENCE: 9301-188-999
; CURRENT APPLICATION NUMBER: US/10/342,887
; CURRENT FILING DATE: 2003-01-15
; PRIOR APPLICATION NUMBER: 60/298,918
; PRIOR FILING DATE: 2001-06-18
; PRIOR APPLICATION NUMBER: 60/380,710
; PRIOR FILING DATE: 2002-05-14
; PRIOR APPLICATION NUMBER: 10/172,118
; PRIOR FILING DATE: 2002-06-14
; NUMBER OF SEQ ID NOS: 2699
; SEQ ID NO 981
; LENGTH: 14082
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-342-887-981

Query Match 100.0%; Score 137; DB 12; Length 14082;
Best Local Similarity 100.0%; Pred. No. 1.2e-60;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAAAATACCAATGGACAAGTGTGAAGCTGTGAAGTGTGAAGTGTGT 60
Db 13424 ATTATAAGGAAAAAGAAAAATACCAATGGACAAGTGTGAAGCTGTGAAGTGTGT 13365

QY 61 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGTGAAGTGTGAAGTGTGAAGTGT 120
Db 13364 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGTGAAGTGTGAAGTGTGT 13305

QY 121 TTTGATGTTAATTAATT 137
Db 13304 TTTGATGTTAATTAATT 13288

RESULT 15
US-10-341-434-108/c
; Sequence 108, Application US/10341434
; Publication No. US20030215835A1
; GENERAL INFORMATION:
; APPLICANT: Origene Technologies
; TITLE OF INVENTION: Differentially Regulated Prostate Cancer Genes
; FILE REFERENCE: 90 204 205 R1

; CURRENT APPLICATION NUMBER: US/10/341,434
; CURRENT FILING DATE: 2003-07-18
; PRIOR APPLICATION NUMBER: US 60/348,164
; PRIOR FILING DATE: 2002-01-15
; PRIOR APPLICATION NUMBER: US 60/348,119
; PRIOR FILING DATE: 2002-01-15
; NUMBER OF SEQ ID NOS: 238
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 108
; LENGTH: 14082
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (703)..(11388)
; OTHER INFORMATION:
US-10-341-434-108

Query Match 100.0%; Score 137; DB 15; Length 14082;
Best Local Similarity 100.0%; Pred. No. 1.2e-60;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 13424 ATTATAAGGAAAAAGAAAAATACCAATGGACAAGTGTGAAGCTGTGAAGTGTGT 13365

QY 61 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGTGAAGTGTGAAGTGTGAAGTGT 120
Db 13364 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGTGAAGTGTGAAGTGTGT 13305

QY 121 TTTGATGTTAATTAATT 137
Db 13304 TTTGATGTTAATTAATT 13288

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Job time : 63.7891 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 14:14:32 ; Search time 15.1147 Seconds
(without alignments)
5030.075 Million cell updates/sec

Title: US-09-966-264D-1

Perfect score: 137

Sequence: 1 attataaaggaaagaaaa.....gtgtttgtgtaattaatt 137

Scoring table: OLIGO_NUC

Gapop 60.0 , Gapext 60.0

Searched: 682709 seqs, 277475446 residues

Word size : 0

Total number of hits satisfying chosen parameters: 1365418

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : Issued Patents NA.*

- 1: /cgn2_6/prodata/2/ina/5A-COMB.seq.*
- 2: /cgn2_6/prodata/2/ina/5B-COMB.seq.*
- 3: /cgn2_6/prodata/2/ina/6A-COMB.seq.*
- 4: /cgn2_6/prodata/2/ina/6B-COMB.seq.*
- 5: /cgn2_6/prodata/2/ina/PCtUS-COMB.seq.*
- 6: /cgn2_6/prodata/2/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	117	85.4	13977	4	US-09-484-970B-60
C 2	42	30.7	19307	3	US-08-836-022A-10
C 3	42	30.7	19307	3	US-09-427-048A-10
C 4	19	13.9	4766	5	PCR-US93-07261-10
C 5	18	13.1	750	4	US-09-107-532A-866
C 6	18	13.1	889	2	US-08-935-886-7
C 7	18	13.1	889	2	US-08-935-886-13
C 8	18	13.1	2076	4	US-09-134-001C-1838
C 9	17	12.4	92	1	US-07-839-751-1
C 10	17	12.4	92	1	US-08-239-427A-1
C 11	17	12.4	92	2	US-08-467-975-1
C 12	17	12.4	92	3	US-08-401-355-1
C 13	17	12.4	1260	3	US-09-081-689-3
C 14	17	12.4	1297	4	US-08-858-207A-66
C 15	17	12.4	1329	4	US-09-305-984-13
C 16	17	12.4	1329	4	US-09-073-541A-13
C 17	17	12.4	1329	4	US-09-493-940-13
C 18	17	12.4	1526	3	US-09-081-689-1
C 19	17	12.4	2817	5	PCR-US93-05944-1
C 20	17	12.4	3356	3	US-09-379-523-4
C 21	17	12.4	3808	2	US-08-417-210A-79
C 22	17	12.4	3808	4	US-09-136-159A-79
C 23	17	12.4	4358	4	US-09-308-453-1
C 24	17	12.4	5152	4	US-10-204-708-48
C 25	17	12.4	5815	4	US-10-204-708-77
C 26	17	12.4	7816	3	US-08-815-808-6
C 27	17	12.4	7797	2	US-08-816-155B-7

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28      17      12.4      7797      3      US-09-079-587-7      Sequence 7, Appli
C 29      17      12.4      8900      4      US-09-305-984-25      Sequence 25, Appli
C 30      17      12.4      8900      4      US-09-073-541A-25      Sequence 25, Appli
C 31      17      12.4      8900      4      US-09-493-940-25      Sequence 25, Appli
C 32      17      12.4      10962      2      US-08-816-155B-6      Sequence 6, Appli
C 33      17      12.4      10962      3      US-09-079-587-6      Sequence 6, Appli
C 34      17      12.4      19877      3      US-08-816-155B-8      Sequence 8, Appli
C 35      17      12.4      19877      3      US-09-079-587-8      Sequence 8, Appli
C 36      17      12.4      28882      4      US-08-361-527-140      Sequence 140, App
C 37      16      11.7      312      4      US-09-343-681A-1844      Sequence 1844, App
C 38      16      11.7      402      4      US-09-621-976-113      Sequence 113, App
C 39      16      11.7      439      4      US-09-397-787-294      Sequence 294, App
C 40      16      11.7      661      4      US-09-702-705-219      Sequence 219, App
C 41      16      11.7      661      4      US-09-736-457-219      Sequence 219, App
C 42      16      11.7      661      4      US-09-614-124B-219      Sequence 219, App
C 43      16      11.7      681      4      US-08-871-325-219      Sequence 219, App
C 44      16      11.7      681      4      US-09-589-184-219      Sequence 219, App
C 45      16      11.7      1247      4      US-09-740-035-1      Sequence 1, Appli

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ALIGNMENTS

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RESULT 1
US-09-484-970B-60/c
; Sequence 60, Application US/09484970B
; Patent No. 6426186
; GENERAL INFORMATION:
; APPLICANT: Jones, Karen A.
; APPLICANT: Volkmut, Wayne G.
; TITLE OF INVENTION: BONE REMODELING GENES
; FILE REFERENCE: PB-0014 US
; CURRENT APPLICATION NUMBER: US/09/484,970B
; CURRENT FILING DATE: 2000-01-18
; NUMBER OF SEQ ID NOS: 172
; SOFTWARE: PERL Program
; SEQ ID NO 60
; LENGTH: 13977
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: Incyte ID No. 6426186 229357.11CBI
; NAME/KEY: unsure
; LOCATION: 11721-11761, 12294, 13969
; OTHER INFORMATION: a, t, c, g, or other
US-09-484-970B-60

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Query Match      85.4%      Score 117; DB 4; Length 13977;
Best Local Similarity 100.0%; Pred. No. 3.6e-51; Mismatches 0; Indels 0; Gaps 0;
Matches 117; Conservative 0;

QY 1 ATTATAAGGAAAGAAATAACGCAATGCGAAGTGTGAAGCTGTGAACCTCAGGTGT 60
DB 13306 ATTATAAGGAAAGAAATAACGCAATGCGAAGTGTGAAGCTGTGAACCTCAGGTGT 13247

QY 61 GCACAAATATCAGGAACACCCGAAACCAAGTAGGATAGAAATAGCATGAGAGCC 117
DB 13246 GCACAAATATCAGGAACACCCGAAACCAAGTAGGATAGAAATAGCATGAGAGCC 13190

```

```

RESULT 2
US-08-836-022A-10
; Sequence 10, Application US/08836022A
; Patent No. 6001557
; GENERAL INFORMATION:
; APPLICANT: Trustees of the University of Pennsylvania
; APPLICANT: Wilson, James M.
; APPLICANT: Fisher, Krishna J.
; APPLICANT: Cher, Shu-Jen
; APPLICANT: Weitzman, Matthew
; TITLE OF INVENTION: Improved Adenovirus Virus and

```

```

CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,022
FILING DATE: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Bak, Mary E.
REGISTRATION NUMBER: 31,215
REFERENCE/DOCKET NUMBER: GNVFN.008PCT
TELECOMMUNICATION INFORMATION:
TELEPHONE: 215-540-9200
TELEFAX: 215-540-5818
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 19307 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: unknown
MOLECULE TYPE: cDNA
SEQUENCE DESCRIPTION: SEQ ID NO: 10:
US-09-427-048A-10

Query Match          30.7%; Score 42; DB 3; Length 19307;
Best Local Similarity 100.0%; Pred.No. 1.5e-12;
Matches 42; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      75  AATCCCAACCAAGTAGGTAGAGTAATAGCATGAGAGC 116
      |||
DB      1563 AACCCCAACCAAGTAGGTAGAGTAATAGCATGAGAGC 1604

RESULT 4
PCT-US93-07261-10
; Sequence 10, Application PC/TUS9307261
; GENERAL INFORMATION:
; TITLE OF INVENTION: PFEMP3 MALARIA ANTIGEN, ANALOGS, ANTIBODIES AND USES THEREOF
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: John H. C. Blasdale
; STREET: One Giralda Farms
; CITY: Madison
; STATE: New Jersey
; COUNTRY: USA
; ZIP: 07940-1000
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: Apple Macintosh
; OPERATING SYSTEM: Macintosh 6.0.5
; SOFTWARE: Microsoft Word 5.1a
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US93/07261
; FILING DATE: 19930805
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/927,531
; FILING DATE: 07-AUG-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Blasdale, John H. C.
; REGISTRATION NUMBER: 31,895
; REFERENCE/DOCKET NUMBER: DX0288K
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 201-822-7398
; TELEFAX: 201-822-7039
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 4766 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA to mRNA
; ORIGINAL SOURCE:
; ORGANISM: Plasmodium falciparum
; STRAIN: Malayan Camp
; IMMEDIATE SOURCE:
; CLONE: p2b1/p12-1

```

FEATURE:
NAME/KEY: CDS
LOCATION: 3..4766
PCT-US93-07261-10

Query Match 13.9%; Score 19; DB 5; Length 4766;
Best Local Similarity 100.0%; Pred. No. 1.1;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 TAAAGCAAAAGAAATAA 23
|||||
DB 190 TAAAGCAAAAGAAATAA 208
|||||

RESULT 5
US-09-107-532A-866
Sequence 866, Application US/09107532A
Patent No. 6593275
GENERAL INFORMATION:
APPLICANT: Lynn A Doucette-Stamm and David Bush
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
NUMBER OF SEQUENCES: 7310
CORRESPONDENCE ADDRESS:
ADDRESSEE: GENOME THERAPEUTICS CORPORATION
STREET: 100 Beaver Street
CITY: Waltham
STATE: Massachusetts
COUNTRY: USA
ZIP: 02154

COMPUTER READABLE FORM:
MEDIUM TYPE: CD-ROM ISO9660
COMPUTER: PC
OPERATING SYSTEM: <Unknown>
SOFTWARE: ASCII
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/107,532A
FILING DATE: 30-Jun-1998
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/085,598
FILING DATE: 14 May 1998
APPLICATION NUMBER: 60/051571
FILING DATE: July 2, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Ariniello, Pamela Deneka
REGISTRATION NUMBER: 40,489
REFERENCE/DOCKET NUMBER: GTC-012
TELEPHONE: (781)893-5007
TELEFAX: (781)893-8277
INFORMATION FOR SEQ ID NO: 866:
SEQUENCE CHARACTERISTICS:
LENGTH: 750 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: circular
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Enterococcus faecium
FEATURE:
NAME/KEY: misc feature
LOCATION: (B) LOCATION 1...750
SEQUENCE DESCRIPTION: SEQ ID NO: 866:
US-09-107-532A-866

Query Match 13.1%; Score 18; DB 4; Length 750;
Best Local Similarity 100.0%; Pred. No. 3.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 CAATGGACAAGTGGTAA 43
|||||

Db 117 CAATGGACAAGTGGTAA 134

RESULT 6
US-08-935-886-7
Sequence 7, Application US/08935886
Patent No. 5994625
GENERAL INFORMATION:
APPLICANT: Melchers, Leo S.
APPLICANT: Sela-Buuriage, Marianne B.
APPLICANT: Bres-Vloemans, Alexandra A.
APPLICANT: Ponstein, Anne S.
APPLICANT: Apotheker-de Groot, Marion
APPLICANT: Cornelissen, Bernardus J. C.
TITLE OF INVENTION: Antifungal Chitin Binding Proteins and DNA
TITLE OF INVENTION: Coding Therefor
NUMBER OF SEQUENCES: 13
CORRESPONDENCE ADDRESS:
ADDRESSEE: Ladass & Parry
STREET: 26 West 61st Street
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10023

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 14.4 MB storage
COMPUTER: IBM PC/XT/AT or compatibles
OPERATING SYSTEM: DOS 6.20
SOFTWARE: Wordperfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/935,886
FILING DATE:
CLASSIFICATION: 800
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/411,640
FILING DATE: 05-APR-1995
APPLICATION NUMBER: PCT/EP93/02790
FILING DATE: 05-OCT-1993
ATTORNEY/AGENT INFORMATION:
NAME: Mass, Clifford J.
REGISTRATION NUMBER: 30,086
REFERENCE/DOCKET NUMBER: U-010139-3
TELEPHONE: (212) 708-1800
TELEFAX: (212) 246-8959
TELEX: 233288
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 889 Base Pairs
TYPE: Nucleic Acid
STRANDEDNESS: Single
TOPOLOGY: Linear
MOLECULE TYPE: cDNA to mRNA
HYPOTHETICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Nicotiana tabacum
STRAIN: Samsun NN
DEVELOPMENTAL STAGE: Mature
TISSUE TYPE: Leaf, wounded
IMMEDIATE SOURCE:
LIBRARY: lambda zap
CLONE: CBP4.4
FEATURE:
NAME/KEY: CDS
LOCATION: 16..639
FEATURE:
NAME/KEY: misc feature
LOCATION: 1..14
FEATURE:
NAME/KEY: misc feature
LOCATION: 866..889
OTHER INFORMATION: /function= "XhoI-linker"

US-08-935-886-7

Query Match 13.1%; Score 18; DB 2; Length 889;
Best Local Similarity 100.0%; Pred. No. 3.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 AAAGGAAAAGAAATAA 23
DB 707 AAAGGAAAAGAAATAA 724

RESULT 7

US-08-935-886-13
; Sequence 13, Application US/08935886
; Patent No. 5994625
; GENERAL INFORMATION:
; APPLICANT: Melchers, Leo S.
; APPLICANT: Sela-Buurlage, Marianne B.
; APPLICANT: Bres-Vloemans, Alexandra A.
; APPLICANT: Ponstein, Anne S.
; APPLICANT: Apotheke-de Groot, Marion
; APPLICANT: Cornelissen, Bernardus J. C.
; TITLE OF INVENTION: Antifungal Chitin Binding Proteins and DNA
; TITLE OF INVENTION: Coding Therefor
; NUMBER OF SEQUENCES: 13
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Ladas & Parry
; STREET: 26 West 61st Street
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10023

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 14.4 MB storage
COMPUTER: IBM PC/XT/AT or compatibles
OPERATING SYSTEM: DOS 6.20
SOFTWARE: Wordperfect 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/935,886
FILING DATE:
CLASSIFICATION: 800
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/411,640
FILING DATE: 05-APR-1995
APPLICATION NUMBER: PCT/EP93/02790
FILING DATE: 05-OCT-1993
ATTORNEY/AGENT INFORMATION:
NAME: Mass, Clifford J.
REGISTRATION NUMBER: 30,086
REFERENCE/DOCKET NUMBER: U-010139-3
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 708-1800
TELEFAX: (212) 246-8959
TELEX: 233288

INFORMATION FOR SEQ ID NO: 13:
SEQUENCE CHARACTERISTICS:
LENGTH: 889 Base Pairs
TYPE: Nucleic Acid
STRANDEDNESS: double
TOPOLOGY: Linear
MOLECULE TYPE: cDNA to mRNA
HYPOTHETICAL: YES
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Nicotiana tabacum
STRAIN: Samsun NN
TISSUE TYPE: Leaf, wounded
IMMEDIATE SOURCE:
CLONE: CP4.4T
US-08-935-886-13

Query Match 13.1%; Score 18; DB 2; Length 889;
Best Local Similarity 100.0%; Pred. No. 3.4;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 AAAGGAAAAGAAATAA 23
DB 726 AAAGGAAAAGAAATAA 743

RESULT 8

US-09-134-001C-1838
; Sequence 1838, Application US/09134001C
; Patent No. 8380370
; GENERAL INFORMATION:
; APPLICANT: Lynn Doucette-Stamm et al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO STAPHYLOCOCCUS
; TITLE OF INVENTION: EPIDERMIDIS FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: GTC-007
; CURRENT APPLICATION NUMBER: US/09/134,001C
; CURRENT FILING DATE: 1998-08-13
; PRIOR APPLICATION NUMBER: US 60/084,964
; PRIOR FILING DATE: 1997-11-08
; PRIOR APPLICATION NUMBER: US 60/055,779
; PRIOR FILING DATE: 1997-08-14
; NUMBER OF SEQ ID NOS: 5674
; SEQ ID NO 1838
; LENGTH: 2076
; TYPE: DNA
; ORGANISM: Staphylococcus epidermidis
US-09-134-001C-1838

Query Match 13.1%; Score 18; DB 4; Length 2076;
Best Local Similarity 100.0%; Pred. No. 3.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TTATAAGGAAAAGAAA 19
DB 1374 TTATAAGGAAAAGAAA 1391

RESULT 9

US-07-839-751-1/c
; Sequence 1, Application US/07839751
; Patent No. 5439809
; GENERAL INFORMATION:
; APPLICANT: Haynes, Joel
; APPLICANT: Klein, Michel H
; TITLE OF INVENTION: Production of Genetically-Engineered
; TITLE OF INVENTION: Vaccines for Aids and Retroviral Diseases
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Sim & McBurney
; STREET: Suite 701, 330 University Avenue
; CITY: Toronto
; STATE: Ontario
; COUNTRY: Canada
; ZIP: M5G 1R7
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA: US/07/839,751
APPLICATION NUMBER: US/07/839,751
FILING DATE: 19920615
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Stewart, Michael I
REGISTRATION NUMBER: 24,973
REFERENCE/DOCKET NUMBER: 1038-267 MLS:JC
TELECOMMUNICATION INFORMATION:
TELEPHONE: (416) 595-1155
TELEFAX: (416) 595-1163
TELEX: 065-24567 SIMBAS
INFORMATION FOR SEQ ID NO: 1:

; SEQUENCE CHARACTERISTICS:

LENGTH: 92 base pairs
 TYPE: NUCLEIC ACID
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)
 US-07-839-751-1

Query Match 12.4%; Score 17; DB 1; Length 92;
 Best Local Similarity 100.0%; Pred. No. 11;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 AGGAAAAAGAAATAAC 24
 |||||

Db 73 AGGAAAAAGAAATAAC 57
 |||||

RESULT 10

US-08-239-427A-1/c
 ; Sequence 1, Application US/08239427A
 ; Patent No. 5571712

; GENERAL INFORMATION:

APPLICANT: Haynes, Joel
 APPLICANT: Klein, Michel H
 APPLICANT: Rovinski, Benjamin
 APPLICANT: Cao, Shi Xian
 TITLE OF INVENTION: Production of Genetically-Engineered
 TITLE OF INVENTION: Vaccines for Aids and Retroviral Diseases
 NUMBER OF SEQUENCES: 2
 CORRESPONDENCE ADDRESS:

ADDRESSEE: Sim & McBurney
 STREET: Suite 701, 330 University Avenue
 CITY: Toronto
 STATE: Ontario
 COUNTRY: Canada
 ZIP: M5G 1R7

; COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patent In Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/239,427A
 FILING DATE: 06-MAY-1994

; CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 839,751

FILING DATE: 12-OCT-1990

ATTORNEY/AGENT INFORMATION:

NAME: Stewart, Michael I

REGISTRATION NUMBER: 24,973

REFERENCE/DOCKET NUMBER: 1038-332 MIS:jb

TELEPHONE: (416) 595-1155

TELEFAX: (416) 595-1163

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 92 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

US-08-239-427A-1

Query Match 12.4%; Score 17; DB 1; Length 92;
 Best Local Similarity 100.0%; Pred. No. 11;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 AGGAAAAAGAAATAAC 24
 |||||

Db 73 AGGAAAAAGAAATAAC 57
 |||||

RESULT 11

US-08-467-975-1/c
 ; Sequence 1, Application US/08467975
 ; Patent No. 5985641

; GENERAL INFORMATION:

APPLICANT: Haynes, Joel
 APPLICANT: Klein, Michel H
 APPLICANT: Rovinski, Benjamin
 APPLICANT: Cao, Shi Xian
 TITLE OF INVENTION: Production of Genetically-Engineered
 TITLE OF INVENTION: Vaccines for Aids and Other Retroviral Diseases
 NUMBER OF SEQUENCES: 2
 CORRESPONDENCE ADDRESS:

ADDRESSEE: Sim & McBurney
 STREET: Suite 701, 330 University Avenue
 CITY: Toronto
 STATE: Ontario
 COUNTRY: Canada
 ZIP: M5G 1R7

; COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patent In Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/467,975
 FILING DATE: 06-JUN-1995

; CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/239,427

FILING DATE: 06-MAY-1994

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07/839,751

FILING DATE: 12-OCT-1990

PRIOR APPLICATION DATA:

APPLICATION NUMBER: GB 8912123.7

FILING DATE: 13-OCT-1989

ATTORNEY/AGENT INFORMATION:

NAME: Stewart, Michael I

REGISTRATION NUMBER: 24,973

REFERENCE/DOCKET NUMBER: 1038-456 MIS:vg

TELEPHONE: (416) 595-1155

TELEFAX: (416) 595-1163

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 92 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

US-08-467-975-1

Query Match 12.4%; Score 17; DB 2; Length 92;
 Best Local Similarity 100.0%; Pred. No. 11;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 AGGAAAAAGAAATAAC 24
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Db 73 AGGAAAAAGAAATAAC 57
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RESULT 12

US-08-401-355-1/c
 ; Sequence 1, Application US/08401355
 ; Patent No. 6291227

; GENERAL INFORMATION:

APPLICANT: Haynes, Joel
 APPLICANT: Klein, Michel H
 TITLE OF INVENTION: Production of Genetically-Engineered
 TITLE OF INVENTION: Vaccines for Aids and Retroviral Diseases
 NUMBER OF SEQUENCES: 2

;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: Sim & McBurney
;; STREET: Suite 701, 330 University Avenue
;; CITY: Toronto
;; STATE: Ontario
;; COUNTRY: Canada
;; ZIP: M5R 1R7
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: PC-DOS/MS-DOS
;; SOFTWARE: Patent In Release #1.0, Version #1.30
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/401,355
;; FILING DATE:
;; CLASSIFICATION: 424
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US 07/839,751
;; FILING DATE: 15-JUN-1992
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Stewart, Michael I
;; REGISTRATION NUMBER: 24,973
;; REFERENCE/DOCKET NUMBER: 1038-424 MIS:as
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (416) 595-1155
;; TELEFAX: (416) 595-1163
;; TELEX: 065-24567 SIMBAS
;; INFORMATION FOR SEQ ID NO: 1:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 92 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: double
;; TOPOLOGY: linear
;; MOLECULE TYPE: DNA (genomic)
;; US-08-401-355-1

Query Match 12.4%; Score 17; DB 3; Length 92;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 AGGAAAAGAAATAAC 24
DB 73 AGGAAAAGAAATAAC 57

RESULT 13
US-09-081-689-3/c
; Sequence 3, Application US/09081689
; Patent No. 6165992
; GENERAL INFORMATION:
; APPLICANT: Wallis, Nicola G.
; APPLICANT: Zalacain, Magdalenaa
; APPLICANT: Throup, John
; APPLICANT: Biswas, Sanjoy
; TITLE OF INVENTION: Histidine Kinase
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Dechert, Price & Rhoads
; STREET: 4000 Bell Atlantic Tower, 1717 Arch Stre
; CITY: Philadelphia
; STATE: PA
; COUNTRY: USA
; ZIP: 19103-2793
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/081,689
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:

;; APPLICATION NUMBER: 60/048,347
;; FILING DATE: 30-MAY-1997
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Falk, Stephen T
;; REGISTRATION NUMBER: 36,795
;; REFERENCE/DOCKET NUMBER: GM10009
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 215-994-2488
;; TELEFAX: 215-994-2222
;; TELEX:
;; INFORMATION FOR SEQ ID NO: 3:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 1260 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: double
;; TOPOLOGY: linear
;; US-09-081-889-3
Query Match 12.4%; Score 17; DB 3; Length 1260;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 7 AAGGAAAAGAAATAA 23
DB 631 AAGGAAAAGAAATAA 615
RESULT 14
US-08-858-207A-66/c
; Sequence 66, Application US/08858207A
; Patent No. 6348328
; GENERAL INFORMATION:
; APPLICANT: Black, Michael
; APPLICANT: Hodgson, John
; APPLICANT: Knowles, David
; APPLICANT: Nicholas, Richard
; APPLICANT: Stodola, Robert
; TITLE OF INVENTION: No. 6348328el Compounds
; NUMBER OF SEQUENCES: 552
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SmithKline Beecham Corporation
; STREET: 709 Swedeland Road
; CITY: King of Prussia
; STATE: PA
; COUNTRY: USA
; ZIP: 19406-0939
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/858,207A
; FILING DATE: 09-MAY-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/017670
; FILING DATE: 14-MAY-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Gimmi, Edward R
; REGISTRATION NUMBER: 38,891
; REFERENCE/DOCKET NUMBER: P50475
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 610-270-4478
; TELEFAX: 610-270-5090
; TELEX:
; INFORMATION FOR SEQ ID NO: 66:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1297 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; US-08-858-207A-66

Search completed: April 6, 2004, 17:42:27
Job time : 16.6147 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 11:42:46 ; Search time 64.9329 Seconds
(without alignments)
8963.144 Million cell updates/sec

Title: US-09-966-264D-1
Perfect score: 137
Sequence: 1 attataaggaagaaagaaa.....gtgttgatgtaattaatt 137

Scoring table: OLIGO NUC
Gapop 60.0 , Gapext 60.0

Searched: 337863 seqs, 2124099041 residues

Word size : 0

Total number of hits satisfying chosen parameters: 6747726

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : N Geneseq 29Jan04: *
1: geneseqn1980s: *
2: geneseqn1990s: *
3: geneseqn2000s: *
4: geneseqn2001as: *
5: geneseqn2001bs: *
6: geneseqn2002s: *
7: geneseqn2003as: *
8: geneseqn2003bs: *
9: geneseqn2003cs: *
10: geneseqn2004s: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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2	137	100.0	158	6	ABK86469 Human Apo
3	137	100.0	200	6	ABK86468 Human dys
4	137	100.0	455	9	AD32504 Human mit
5	137	100.0	996	6	ABK86497 Human Apo
6	137	100.0	1230	6	ABK86462 Human Apo
7	137	100.0	1234	6	ABK86463 Human Apo
8	137	100.0	2691	6	ABK81996 Human dys
9	137	100.0	8689	6	ABK82000 Human dys
10	137	100.0	11443	6	ABK82002 DNA encod
11	137	100.0	13957	6	ABK81959 CDNA enco
12	137	100.0	13957	6	ABK10904 Human bre
13	137	100.0	13957	6	ABN95786 Gene #228
14	137	100.0	13957	6	ABK86900 Human dys
15	117	85.4	13977	6	ABK70403 Human bon
16	46	33.6	959	6	ABN74601 Bovine em
17	42	30.7	13815	2	AAV18885 Mus muscu
18	42	30.7	13815	6	ABK81960 CDNA enco
19	42	30.7	13815	6	ABK81960 CDNA enco
20	42	30.7	13815	6	ABK81960 CDNA enco
21	41	29.9	60	6	ABK86471 Human apo
22	40	29.2	108	6	ABK86467 Human apo
23	22	16.1	22	6	ABK86479 Human apo

c	24	20	14.6	690	6	ABZ08220 Human leu
c	25	19	13.9	716	4	AAK64599 Human inn
c	26	19	13.9	4766	2	AAQ70102 Malarial
c	27	19	13.9	7326	3	AAA70095 Plasmodiu
c	28	19	13.9	42521	2	AAZ32013 Human MFT
c	29	19	13.9	42521	5	AAC90070 U64857 cD
c	30	18	13.1	18	6	ABK86487 Human apo
c	31	18	13.1	324	4	AAZ33883 Tobacco p
c	32	18	13.1	407	4	AAK75817 Human inn
c	33	18	13.1	408	4	AAK75816 Human inn
c	34	18	13.1	476	9	ADB51265 Primary r
c	35	18	13.1	492	7	ABZ76803 Nicotiana
c	36	18	13.1	569	3	AAC55084 Arabidops
c	37	18	13.1	717	7	ACA33643 Prokaryot
c	38	18	13.1	750	9	ABN98770 Arabidops
c	39	18	13.1	878	6	AAQ62459 Chitin bi
c	40	18	13.1	889	2	AAQ62455 Chitin bi
c	41	18	13.1	896	3	AAC37805 Arabidops
c	42	18	13.1	1028	3	AAC37406 Arabidops
c	43	18	13.1	1155	3	AAC36681 Arabidops
c	44	18	13.1	2076	6	ABN92375 Staphyloc
c	45	18	13.1	2076	6	ABN92375 Staphyloc

ALIGNMENTS

RESULT 1
ID ABK86496 standard; DNA; 137 BP.
XX AC ABK86496;
XX DT 27-AUG-2002 (first entry)
XX DE Human Apo-dystrophin-4 inversion sequence.
XX KW Human; ds; apo-dystrophin-4; inversion sequence; gene therapy;
XX KW protein truncation; muscular dystrophy; leukaemia.
XX OS Homo sapiens.
XX FH Key Location/Qualifiers
FT CDS 3..137
FT FT /*tag= a
FT FT /product= "Apo-dystrophin-4 peptide appearing as
FT FT AAU98738"
FT FT /partial
FT FT /note= "No start or stop codon shown"
FT FT /transl_except= (pos:21..23,aa:Xaa)
FT FT /transl_except= (pos:48..50,aa:Xaa)
FT FT /transl_except= (pos:93..95,aa:Xaa)
FT FT /transl_except= (pos:123..125,aa:Xaa)
FT FT /transl_except= (pos:129..131,aa:Xaa)
FT FT /note= "Xaa= unknown, encoded by in frame stop codon"

GB2368064-A.

24-APR-2002.

16-JAN-2001; 2001GB-00001124.

30-SEP-2000; 2000US-0237079P.

(IMCR) IMPERIAL CANCER RES TECHNOLOGY LTD.
(BARB/) BARBER E.

Barber E;

WPI; 2002-425042/46.

P-PSDB; AAU98737.

New human regulatory polynucleotide, useful for treating disorders

PT associated with protein truncation, particularly muscular dystrophy, and
PT related peptides and antibodies.
PS Claim 1; Page 169; 222pp; English.
XX The invention relates to a polynucleotide (I) comprising, or consisting
CC of, apo-dystrophin-4 inversion sequence appearing as ABK86496, or its
CC functional equivalents (e.g. the apo-dystrophin-4 cDNA sequence appearing
CC as ABK86497). Also included are polynucleotides that hybridise to either
CC strand of (I), a vector containing (I), a cell containing (I) or the
CC vector, proteins and peptides encoded by (I), a protein homologous with
CC human dystrophin that is expressed on cell surfaces in vivo antibodies
CC (Ab) specific for the protein and method of screening for leukemia cells
CC by analysing DNA for presence of (I) or by detecting presence of (II).
CC The apo-dystrophin-4 inversion sequence is a regulatory element that
CC controls expression (transcription and translation) of associated DNA,
CC and may allow read-through of stop codons. The apo-dystrophin-4 inversion
CC sequence is used in gene therapy of diseases associated with truncation
CC of proteins, particularly muscular dystrophy and also leukaemia, but more
CC generally (I) is a regulatory sequence used to control expression of any
CC attached gene. Analysis of DNA for (I), or detection of proteins (II)
CC encoded by (I), can be used to screen for leukaemic cells and related
CC diseases. Antibodies raised against (II) can be used therapeutically, to
CC inhibit (II) activity, also to detect (II) in screening assays. The
CC present sequence is the apo-dystrophin-4 inversion sequence and upstream
CC genomic region
XX
SQ Sequence 137 BP; 56 A; 19 C; 32 G; 30 T; 0 U; 0 Other;
Query Match 100.0%; Score 137; DB 6; Length 137;
Best Local Similarity 100.0%; Pred. No. 4.8e-59;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAGAAATAACGCAATGCAAGTGGTGAAGCTGTGAAGCTCAGGTGT 60
DB 1 ATTATAAGGAAAGAAATAACGCAATGCAAGTGGTGAAGCTGTGAAGCTCAGGTGT 60
QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGAATAGCATGAGAGCCGTG 120
DB 61 GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGAATAGCATGAGAGCCGTG 120
QY 121 TTTGATGTTAATTAATT 137
DB 121 TTTGATGTTAATTAATT 137
RESULT 2
ABK86469/c
ID ABK86469 standard; cDNA; 158 BP.
XX AC ABK86469;
XX 27-AUG-2002 (first entry)
XX Human apo-dystrophin-4 cDNA fragment with inversion breakpoint #2.
XX Human; ss; apo-dystrophin-4; inversion sequence; gene therapy;
XX protein truncation; muscular dystrophy; leukaemia; dystrophin.
XX Homo sapiens.
XX Key Location/Qualifiers
XX misc_recomb 137
XX /tag= a
XX /label= Inversion_breakpoint
XX GB2368064-A.
XX 24-APR-2002.
XX 16-JAN-2001; 2001GB-00001124.
XX 30-SEP-2000; 2000US-0237079P.
PR

XX (TMCR) IMPERIAL CANCER RES TECHNOLOGY LTD.
PA (BARE//) BARBER E.
XX Barber E;
XX WPI; 2002-429042/46.
XX New human regulatory polynucleotide, useful for treating disorders
PT associated with protein truncation, particularly muscular dystrophy, and
PT related peptides and antibodies.
XX Disclosure; Fig 16B; 222pp; English.
XX The invention relates to a polynucleotide (I) comprising, or consisting
CC of, apo-dystrophin-4 inversion sequence appearing as ABK86496, or its
CC functional equivalents (e.g. the apo-dystrophin-4 cDNA sequence appearing
CC as ABK86497). Also included are polynucleotides that hybridise to either
CC strand of (I), a vector containing (I), a cell containing (I) or the
CC vector, proteins and peptides encoded by (I), a protein homologous with
CC human dystrophin that is expressed on cell surfaces in vivo antibodies
CC (Ab) specific for the protein and method of screening for leukemia cells
CC by analysing DNA for presence of (I) or by detecting presence of (II).
CC The apo-dystrophin-4 inversion sequence is a regulatory element that
CC controls expression (transcription and translation) of associated DNA,
CC and may allow read-through of stop codons. The apo-dystrophin-4 inversion
CC sequence is used in gene therapy of diseases associated with truncation
CC of proteins, particularly muscular dystrophy and also leukaemia, but more
CC generally (I) is a regulatory sequence used to control expression of any
CC attached gene. Analysis of DNA for (I), or detection of proteins (II)
CC encoded by (I), can be used to screen for leukaemic cells and related
CC diseases. Antibodies raised against (II) can be used therapeutically, to
CC inhibit (II) activity, also to detect (II) in screening assays. The
CC present sequence is a apo-dystrophin-4 cDNA fragment showing an inversion
CC breakpoint (recombination signal sequence) similar to that in with human
CC dystrophin
XX
SQ Sequence 158 BP; 35 A; 35 C; 20 G; 68 T; 0 U; 0 Other;
Query Match 100.0%; Score 137; DB 6; Length 158;
Best Local Similarity 100.0%; Pred. No. 4.8e-59;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATTATAAGGAAAGAAATAACGCAATGCAAGTGGTGAAGCTGTGAAGCTCAGGTGT 60
DB 137 ATTATAAGGAAAGAAATAACGCAATGCAAGTGGTGAAGCTGTGAAGCTCAGGTGT 78
QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGAATAGCATGAGAGCCGTG 120
DB 77 GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGAATAGCATGAGAGCCGTG 18
QY 121 TTTGATGTTAATTAATT 137
DB 17 TTTGATGTTAATTAATT 1
RESULT 3
ABK86468/c
ID ABK86468 standard; DNA; 200 BP.
XX AC ABK86468;
XX 27-AUG-2002 (first entry)
XX Human dystrophin genomic DNA with inversion breakpoint #2.
XX Human; ds; apo-dystrophin-4; inversion sequence; gene therapy;
XX protein truncation; muscular dystrophy; leukaemia; dystrophin.
XX Homo sapiens.
XX Key Location/Qualifiers
XX misc_recomb 42
FT

```
FT      /*tag= a
FT      /label= Inversion_breakpoint
FT      misc_recomb 179
FT      /*tag= b
FT      /label= Inversion_breakpoint
XX
XX GB2368064-A.
XX
XX 24-APR-2002.
XX
XX 16-JAN-2001; 2001GB-00001124.
XX
XX 30-SEP-2000; 2000US-0237079P.
XX
XX (IMCR ) IMPERIAL CANCER RES TECHNOLOGY LTD.
XX (BARB/) BARBER E.
XX
XX Barber E;
XX
XX WPI; 2002-429042/46.
XX
XX New human regulatory polynucleotide, useful for treating disorders
XX associated with protein truncation, particularly muscular dystrophy, and
XX related peptides and antibodies.
XX
XX Disclosure; Fig 16B; 222pp; English.
XX
XX The invention relates to a polynucleotide (I) comprising, or consisting
XX of, apo-dystrophin-4 inversion sequence appearing as A3K86496, or its
XX functional equivalents (e.g. the apo-dystrophin-4 cDNA sequence appearing
XX as AK86497). Also included are polynucleotides that hybridise to either
XX strand of (I), a vector containing (I), a cell containing (I) or the
XX vector, proteins and peptides encoded by (I), a protein homologous with
XX human dystrophin that is expressed on cell surfaces in vivo antibodies
XX (Ab) specific for the protein and method of screening for leukemia cells
XX by analysing DNA for presence of (I) or by detecting presence of (II).
XX The apo-dystrophin-4 inversion sequence is a regulatory element that
XX controls expression (transcription and translation) of associated DNA,
XX and may allow read-through of stop codons. The apo-dystrophin-4 inversion
XX sequence is used in gene therapy of diseases associated with truncation
XX of proteins, particularly muscular dystrophy and also leukaemia, but more
XX generally (I) is a regulatory sequence used to control expression of any
XX attached gene. Analysis of DNA for (I), or detection of proteins (II)
XX encoded by (I), can be used to screen for leukaemic cells and related
XX diseases. Antibodies raised against (II) can be used therapeutically, to
XX inhibit (II) activity, also to detect (II) in screening assays. The
XX present sequence is a human dystrophin genomic DNA fragment showing
XX inversion breakpoints (recombination signal sequence) similar to that in
XX apo-dystrophin-4
XX
XX Sequence 200 BP; 44 A; 40 C; 28 G; 88 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 137; DB 6; Length 200;
XX Best Local Similarity 100.0%; Pred. No. 4.8e-59;
XX Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1 ATTATAAGGAAAAAGAAATACGCAATGCAAGTGGTGAAGCTGTGCACTCAGGTGT 60
XX
XX 179 ATTATAAGGAAAAAGAAATACGCAATGCAAGTGGTGAAGCTGTGCACTCAGGTGT 120
XX
XX 61 GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGAATACATCAGAACCGGTG 120
XX
XX 119 GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGAATACATCAGAACCGGTG 60
XX
XX 121 TTGTGCTTTAATTAATT 137
XX
XX 59 TTGTGCTTTAATTAATT 43
XX
XX RESULT 4
XX ID ADD32504/c
XX ID ADD32504 standard; DNA; 455 BP.
XX
```

```
AC ADD32504;
XX
XX 15-JAN-2004 (first entry)
XX
XX Human mitochondrial DNA sequence SEQ ID NO:274.
XX
XX ds; human; array; mitochondrial; hybridisation; energy-metabolism;
XX mitochondrial disease; oxidative phosphorylation dysfunction;
XX oxidative stress; apoptosis; aging.
XX
XX Homo sapiens.
XX
XX WO2003020220-A2.
XX
XX 13-MAR-2003.
XX
XX 30-AUG-2002; 2002WO-US027886.
XX
XX 30-AUG-2001; 2001US-0316323P.
XX
XX 31-AUG-2001; 2001CA-02356540.
XX
XX (UYEM-) UNIV EMORY.
XX
XX Wallace DC, Levy S, Kerstann K, Procaccio V;
XX
XX WPI; 2003-300821/29.
XX
XX Array containing probes for genes involved in mitochondrial biology,
XX useful for determining mitochondrial biology gene expression profiles for
XX use in diagnosing pathologies and identifying biochemical pathways.
XX
XX Claim 2; SEQ ID NO 274; 201pp; English.
XX
XX The invention relates to a novel array comprising at least two isolated
XX nucleotide molecules, each molecule having a sequence capable of uniquely
XX hybridising to a nucleic acid molecule which is an expression product of
XX a gene involved in mitochondrial biology. The array comprises two or more
XX isolated nucleic acid molecules or spots, each molecule having a sequence
XX chosen from sequence of 994 human probes and 2046 mouse probes. An array
XX of the invention is useful for determining an expression profile of a
XX mouse or human sample containing nucleic acid, by contacting the array
XX measuring hybridisation of nucleic acid in the sample to the array to
XX produce an expression profile. The array is also useful for determining
XX an expression profile of a first labelled sample containing nucleic acid
XX relative to a second, differently labelled sample containing nucleic
XX acid. The second sample is a reference or a standard. An array is useful
XX for determining an expression profile diagnostic of an energy-metabolism-
XX related physiological condition. An array of the invention is useful for
XX determining mitochondrial biology gene expression profiles of organisms,
XX such as human, mice and closely related species, tissue and organs of
XX such organisms, which are useful for determining expression profiles
XX diagnostic of energy metabolism-related physiological conditions,
XX diagnosing such physiological conditions, identifying biochemical
XX pathways, genes, and mutations involved in such physiological conditions,
XX identifying therapeutic agents useful for preventing and/or treating such
XX physiological conditions, evaluating and/or monitoring the efficacy of
XX such therapies, and creating and identifying animal models of human
XX energy metabolism-related physiological conditions. An array is also
XX useful for defining expression signatures or profiles for mitochondrial
XX diseases, as well as distinguishing clinical disorders that result from
XX oxidative phosphorylation (OXPHOS) dysfunction, oxidative stress,
XX apoptosis and aging. An array of the invention contains probes of genes
XX not previously recognised to participate in mitochondrial biology. The
XX sequences shown in ADD3231-ADD3323 represent human mitochondrial DNA
XX clones used to make the probes of the invention. Some sequences are not
XX present, these are SEQ ID NO's 295, 1174, 1213, 1700, 1728, 1730, 1905,
XX 1906, 2408 and 2643.
XX
XX Sequence 455 BP; 127 A; 80 C; 63 G; 185 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 137; DB 9; Length 455;
XX Best Local Similarity 100.0%; Pred. No. 4.7e-59;
```


QY	1	ATTATAAGGAAAAAGAAATACGCAATGCGACCAAGTGGTGAAGCTGTGAACTCAGGTGT	60
Db	2033	ATTATAAGGAAAAAGAAATACGCAATGCGACCAAGTGGTGAAGCTGTGAACTCAGGTGT	1974
QY	61	GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGAAGTGAAGTGAAGCCGTG	120
Db	1973	GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGAAGTGAAGTGAAGCCGTG	1914
QY	121	TTTGATGTTAATTAAAT	137
Db	1913	TTTGATGTTAATTAAAT	1897
RESULT 9			
ABX82000/c			
ID ID ABK92000 standard; DNA; 8689 BP.			
XX	AC	ABK92000;	
XX	DT	13-AUG-2002 (first entry)	
XX	DE	DNA encoding mini-dystrophin protein deltaH2-R19.	
XX	KW	Mini-dystrophin peptide; spectrin-like repeat domain; muscle disease;	
XX	KW	Duchenne's muscular dystrophy; DMD; dystrophin; ds.	
XX	OS	Homo sapiens.	
XX	OS	Synthetic.	
XX	FN	WO200229056-A2.	
XX	PD	11-APR-2002.	
XX	PF	04-OCT-2001; 2001WO-US031126.	
XX	PR	06-OCT-2000; 2000US-0238848P.	
XX	PA	(UNMI) UNIV MICHIGAN.	
XX	PI	Chamberlain JS, Harper SQ;	
XX	DR	WPI; 2002-435334/46.	
XX	FT	A composition for preparing therapeutic drugs, has a mini-dystrophin	
XX	PT	peptide comprising a specific number of spectrin-like repeat domains, or	
XX	PT	a nucleic acid sequence encoding the mini-dystrophin peptide.	
XX	PS	Disclosure; Fig 15; 145pp; English.	
XX	CC	The invention describes a composition comprising a mini-dystrophin	
XX	CC	peptide comprising a spectrin-like repeat domain, where the domain	
XX	CC	comprises n spectrin-like repeats, and contains no more than n spectrin-	
XX	CC	like repeats, where n is an even number between 4-24, or a nucleic acid	
XX	CC	encoding a mini-dystrophin peptide. The mini-dystrophin peptide or the	
XX	CC	polynucleotide encoding it is useful as a medicament, for preparing a	
XX	CC	drug for therapeutic application and in the preparation of a composition	
XX	CC	for treatment of muscle disease, e.g. Duchenne's muscular dystrophy	
XX	CC	(DMD). This sequence represents a mini-dystrophin sequence of the	
XX	CC	invention	
XX	SQ	Sequence 8689 BP; 2721 A; 1804 C; 1861 G; 2303 T; 0 U; 0 Other;	
Query Match			
Best Local Similarity 100.0%; Score 137; DB 6; Length 8689;			
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0			
QY	1	ATTATAAGGAAAAAGAAATACGCAATGCGACCAAGTGGTGAAGCTGTGAACTCAGGTGT	60
Db	8031	ATTATAAGGAAAAAGAAATACGCAATGCGACCAAGTGGTGAAGCTGTGAACTCAGGTGT	7971
QY	61	GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGAAGTGAAGTGAAGCCGTG	120
Db	7971	GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAAGTGAAGTGAAGTGAAGCCGTG	7971

KW	adenosine deaminase deficiency; severe combined immune deficiency; PAH; beta-chain; haemoglobin gene; beta-thalassaemia; sickle cell disease; low density lipoprotein gene; familial hypercholesterolaemia; hypoxanthine-guanine phosphoribosyltransferase; Lesch-Nyhan syndrome; phenylalanine hydroxylase gene; gene therapy; phenylketonuria; dystrophin gene; muscular dystrophy; cystic fibrosis; immunostimulant; human cystic fibrosis transmembrane conductance regulator gene; antianemic; antilipemic; nontropic; cytostatic; dermatological; human; alpha-1-antitrypsin; lysosomal glucocerebrosidase; ADA; HPRT; lysosomal aryl-sulfatase A; omithine transcarbamylase; ARSA; OTC; NP; purin nucleoside phosphorylase; gene; ds.
XX	Homo sapiens.
OS	US2002102731-A1.
XX	XX
PN	XX
XX	XX
XX	01-AUG-2002.
PP	XX
XX	XX
PF	12-FEB-2001; 2001US-00782378.
XX	XX
XX	02-OCT-2000; 2000US-0237747P.
PR	XX
XX	XX
XX	(UYNY) UNIV NEW YORK STATE RES FOUND.
PA	Hearing P, Bahou WF, Sandalon Z, Gnatenko DV;
XX	WPI; 2002-690619/74.
XX	XX
DR	XX
XX	Producing vector, by introducing vector having nucleotide sequence, adenovirus inverted terminal repeats and packaging sequence, and adenovirus terminal repeat, into cell, and culturing cell.
PT	XX
PT	XX
PT	XX
XX	XX
XX	XX
PS	Disclosure; Page 122-128; 191pp; English.
XX	XX
CC	The present invention relates to a new method of producing a vector. The method involves introducing recombinant vector having nucleotide sequence (NS) having 5' and 3' end, left and right inverted terminal repeats of adenovirus flanking NS, adenovirus packaging sequence linked to inverted terminal repeat, and adeno-associated virus terminal repeat linked to 3' end of NS, into cell expressing adenovirus early gene lacking from vector ; and culturing cell to produce another vector. The method is useful for generating vectors, especially mad vectors. The method is useful in transferring nucleotide sequences of interest into a cell, for gene transfer applications (e.g. gene therapy) in vitro, ex vivo and in vivo.
CC	The nucleotide sequences are useful for treating diseases associated with it, i.e. adenosine deaminase gene associated with adenosine deaminase deficiency with severe combined immune deficiency, beta-chain of hemoglobin gene associated with beta-thalassaemia and sickle cell disease, receptor for low density lipoprotein gene associated with familial hypercholesterolaemia, hypoxanthine-guanine phosphoribosyltransferase associated with Lesch-Nyhan syndrome, phenylalanine hydroxylase (PAH) gene associated with phenylketonuria, dystrophin gene associated with muscular dystrophy, and human cystic fibrosis transmembrane conductance regulator gene associated with cystic fibrosis. The present nucleic acid sequence represents a human disease gene sequence that was used in the methods of the invention.
CC	XX
XX	XX
SQ	Sequence 13957 BP; 4602 A; 2781 C; 3122 G; 3452 T; 0 U; 0 Other; Query Match 100.0%; Score 137; DB 6; Length 13957; Best Local Similarity 100.0%; Pred. No. 4.6e-59; Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	1 ATTATAAGGAAAAGAAAATATACGCAATGGACAACTGGTGAAGCTGTGAACTCAGGTGT 60
Db	13299 ATTATAAGGAAAAGAAAATATACGCAATGGACAACTGGTGAAGCTGTGAACTCAGGTGT 13240
QY	61 GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAGTGAATAATAGCATGAGAAGCCGTG 120
Db	13239 GCACAAATTATCAGGAACACCCCAAAACCAAAAGTGAGTGAATAATAGCATGAGAAGCCGTG 13180
QY	121 TTTTGATGTTAAATTAAT 137

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 14:12:12 ; Search time 505.195 Seconds
(without alignments)
8098.097 Million cell updates/sec

Title: US-09-966-264d-1

Perfect score: 137

Sequence: 1 attataaagaaagaaagaaaa.....gtgttgatgtaattaatt 137

Scoring table: OLIGO_NUC

Gapop 60.0 , Gapext 60.0

Searched: 27513289 seqs, 14931090276 residues

Word size : 0

Total number of hits satisfying chosen parameters: 55026578

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

EST:*

1: em_estba:*

2: em_esthum:*

3: em_estin:*

4: em_estmu:*

5: em_estov:*

6: em_estpl:*

7: em_estro:*

8: em_hci:*

9: gb_est1:*

10: gb_est2:*

11: gb_hci:*

12: gb_est3:*

13: gb_est4:*

14: gb_est5:*

15: em_estfun:*

16: em_estom:*

17: em_gss_hum:*

18: em_gss_inv:*

19: em_gss_pln:*

20: em_gss_vrt:*

21: em_gss_fun:*

22: em_gss_mam:*

23: em_gss_mus:*

24: em_gss_pro:*

25: em_gss_rod:*

26: em_gss_pbg:*

27: em_gss_vrl:*

28: gb_gss1:*

29: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	137	100.0	373	10	AW580404
C 2	137	100.0	396	14	H89576
C 3	137	100.0	455	9	AA427831
C 4	137	100.0	462	14	CB045405

C 5	137	100.0	473	14	CA389721
C 6	137	100.0	489	28	AQ015185
C 7	137	100.0	647	10	AW664684
C 8	137	100.0	696	10	AW950480
C 9	137	100.0	710	9	AV725574
C 10	137	100.0	727	12	BG567176
C 11	137	100.0	740	14	CD356811
C 12	137	100.0	745	14	CD357556
C 13	137	100.0	1121	12	BM546012
C 14	136	99.3	743	14	CD110642
C 15	120	87.6	352	10	AW385154
C 16	112	81.8	862	14	CB962272
C 17	111	81.0	353	10	AW580423
C 18	88	64.2	911	12	BI752714
C 19	96	62.8	375	10	AW607064
C 20	57	41.6	551	9	AA621932
C 21	54	39.4	270	14	F25920
C 22	54	39.4	841	10	BF791019
C 23	48	35.0	265	14	F18701
C 24	46	33.6	631	14	CB465228
C 25	43	31.4	641	9	AI528613
C 26	42	30.7	268	10	BI178429
C 27	42	30.7	291	10	BI174122
C 28	42	30.7	291	10	BB339107
C 29	42	30.7	296	10	BB041920
C 30	42	30.7	329	10	BB085328
C 31	42	30.7	390	9	AI324317
C 32	42	30.7	607	10	BE370292
C 33	42	30.7	610	9	AA146038
C 34	42	30.7	643	14	CB850037
C 35	42	30.7	735	14	CB570707
C 36	42	30.7	4437	11	AK036936
C 37	40	29.2	734	10	BF674672
C 38	40	29.2	924	13	BF455062
C 39	32	23.4	210	10	BI171938
C 40	32	23.4	418	13	BY377466
C 41	31	22.6	293	10	BB307434
C 42	30	21.9	651	13	BU338637
C 43	30	21.9	733	13	BU357193
C 44	27	19.7	945	10	BF180441
C 45	25	18.2	306	10	BB086855

ALIGNMENTS

RESULT 1
AW580404/c
LOCUS
DEFINITION PM2-HT0451-080100-003-h09 HT0451 Homo sapiens cDNA, mRNA sequence.
ACCESSION AW580404
VERSION AW580404.1 GI:7255453
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 373)
HGCP <http://www.ludwig.org.br/ORESTES>.
The FAPESP/LICR Human Cancer Genome Project
UNPUBLISHED (1999)
CONTACT: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL
(<http://www.ludwig.org.br/scripts/gethtml2.pl?cl=PM2&cl2=PM2-HT0451-080100-003-h09&l3=2000-01-08&t4=1>)


```

/dev_stages="8-9 weeks"
/lab_host="DH10B"
/clone_lib="Soares total fetus NB2HF8 9w"
polylinker: Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
was prepared from mRNA obtained from pooled 8-9 week
(total) fetus material with a Not I - oligo(dT) primer [5'
TGTACCAATCTGAAGTGGAGCGCGCCCTTAATTTTCTTTTCTTTT 3'].
Double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pT73 vector. Library
went through one round of normalization, and was
constructed by Bento Soares and M. Fatima Bonaldo. "

ORIGIN
Query Match      100.0%; Score 137; DB 9; Length 455;
Best Local Similarity 100.0%; Pred. No. 3e-57;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAGAAATTAACCAATGGACAAAGTGTGAAGCTGTGAAGTCAAGTGT 60
DB 155 ATTATAAGGAAAGAAATTAACCAATGGACAAAGTGTGAAGCTGTGAAGTCAAGTGT 96

QY 61 GCACAATTATCAGGACACCCCAAAACCAAGTAGGAGTAAATAGCATGAGAGCCGTTG 120
DB 95 GCACAATTATCAGGACACCCCAAAACCAAGTAGGAGTAAATAGCATGAGAGCCGTTG 36

QY 121 TTGTGATGTTAATTAATT 137
DB 35 TTGTGATGTTAATTAATT 19

RESULT 4
CB045405/c
LOCUS
DEFINITION
NISC_gc10c01.y1 NCI_CGAP_Col7 Homo sapiens cDNA clone IMAGE:3218281
5', mRNA sequence.
CB045405
VERSION
KEYWORDS
SOURCE
Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 462)
AUTHORS
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL
Unpublished (1997)
COMMENT
Contact: Robert Strausberg, Ph.D.
Email: cgapsb@mail.nih.gov
cDNA Library Preparation:
The I.M.A.G.E. Consortium/LLNL
DNA Sequencing by: National Institutes of Health Intramural
Sequencing Center (NISC)
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
info@image.llnl.gov
Plate: LLAM7870 row: E column: 2
Seq primer: M13Rpl reverse primer (ABI).
Location/Qualifiers
1. .462
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:3218281"
/tissue_type="juvenile granulosa tumor"
/lab_host="DH10B"
/clone_lib="NCI CGAP Col7"
/note="Organ: colon; Vector: pCMV-SPORT6; Site_1: SalI;
Site_2: NotI; Cloned unidirectionally. Primer: Oligo dT.
Library constructed by Life Technologies."

ORIGIN
Query Match      100.0%; Score 137; DB 14; Length 462;
Best Local Similarity 100.0%; Pred. No. 3e-57;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAGAAATTAACCAATGGACAAAGTGTGAAGCTGTGAAGTCAAGTGT 60
DB 432 ATTATAAGGAAAGAAATTAACCAATGGACAAAGTGTGAAGCTGTGAAGTCAAGTGT 373

QY 61 GCACAATTATCAGGACACCCCAAAACCAAGTAGGAGTAAATAGCATGAGAGCCGTTG 120
DB 372 GCACAATTATCAGGACACCCCAAAACCAAGTAGGAGTAAATAGCATGAGAGCCGTTG 313

QY 121 TTGTGATGTTAATTAATT 137
DB 312 TTGTGATGTTAATTAATT 296

RESULT 5
CA389721/c
LOCUS
DEFINITION
cs101f03.y3 Human Retinal pigment epithelium/choroid cDNA
(Un-normalized, unamplified): cs Homo sapiens cDNA clone cs101f03
5', mRNA sequence.
CA389721
VERSION
KEYWORDS
SOURCE
Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 473)
AUTHORS
Wistow,G., Bernstein,S.L., Wyatt,M.K., Farris,R.N., Behal,A.,
Touchman,J.W., Bouffard,G., Smith,D. and Peterson,K.
Expressed sequence tag analysis of human RPE/choroid for the
NSIBank Project: Over 6000 non-redundant transcripts, novel genes
and splice variants
Mol. Vis. 8 (4), 205-220 (2002)
JOURNAL
MEDLINE
PUBMED
COMMENT
Contact: Wistow G
Section on Molecular Structure and Function
National Eye Institute
6/331, NIH, Bethesda, MD 20892-2740, USA
Tel: 301 402 3452
Fax: 301 496 0078
Email: graeme@helix.nih.gov
Plate: 101 row: f column: 03
Seq primer: M13Rpl reverse primer (ABI).
Location/Qualifiers
1. .473
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="cs101f03"
/tissue_type="RPE/choroid"
/dev_stage="Adult"
/lab_host="EMDH10B"
/clone_lib="Human Retinal pigment epithelium/choroid cDNA
(Un-normalized, unamplified): cs
/note="Organ: Eye; Vector: pCMVSPORT6; Two different donor
eyes (75-80 years old) yielded approximately 600 mg of
dissected RPE/choroid tissue. This in turn yielded 340 ug
of total RNA and 7 ug of mRNA. A directionally cloned cDNA
library in the pCMVSPORT6 vector was constructed at Life
Technologies (Rockville, MD; now part of Invitrogen Corp),
essentially following the protocols of the SuperScript
Plasmid System (Invitrogen Corp).
<http://www.invitrogen.com/>. The library code
designation was cs. For this library, cDNA inserts were
cloned into the NotI/MluI sites of the vector. EST
analysis was performed on the unamplified library at the
NIH Intramural Sequencing Center (NISC)."

```

Db 298 GCACAAATTATCAGGACACCCCAACCAAGTGAGGTAGAAATACATGAGAACCGTG 239
 QY 121 TTTCATGTTTAATTAATT 137
 Db 238 TTTCATGTTTAATTAATT 222

 RESULT 7
 AW664684/c
 LOCUS
 DEFINITION
 647 bp mRNA linear EST 06-APR-2000
 h184e10.x1 Soares NFL_T_GBC_S1 Homo sapiens cDNA clone
 IMAGE:2979018 3' similar to gb:U18333 DYSTROPHIN (HUMAN);, mRNA
 sequence.
 AW664684
 EST.
 SOURCE
 Homo sapiens (human)
 ORGANISM
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 647)
 NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
 REFERENCE
 AUTHORS
 TITLE
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 JOURNAL
 Unpublished (1997)
 COMMENT
 Contact: Robert Strausberg, Ph.D.
 Email: cgabs-r@mail.nih.gov
 This clone is available royalty-free through LLNL; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 Seq primer: -40UP from Gibco
 High quality sequence stop: 365.
 FEATURES
 Location/Qualifiers
 1..647
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:2979018"
 /lab_host="DH10B"
 /clone_lib="Soares_NFL_T_GBC_S1"
 /notes="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with
 a modified polylinker; Site 1: Not 1; Site 2: Eco RI;
 Equal amounts of plasmid DNA from three normalized
 libraries (fetal lung NbH19W, testis NHT, and B-cell
 NCI CGAP GCBI) were mixed and ss circles were made in
 vitro. Following HAP purification, this DNA was used as
 tracer in a subtractive hybridization reaction. The driver
 was PCR-amplified cDNAs from pools of 5,000 clones made
 from the same 3 libraries. The pools consisted of
 I.M.A.G.E. clones 297480-302087, 682632-687239,
 726408-726711, and 729096-731399. Subtraction by Bento
 Soares and M. Fatima Bonaldo."
 ORIGIN
 Query Match 100.0%; Score 137; DB 10; Length 647;
 Best Local Similarity 100.0%; Pred.No. 3e-57;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ATTATAAGGAAAAGAAAATAACCCCAATGGACAAAGTGGTGAAGCTGTGAAGTGTGT 60
 Db 151 ATTATAAGGAAAAGAAAATAACCCCAATGGACAAAGTGGTGAAGCTGTGAAGTGT 92
 QY 61 GCACAAATTATCAGGAACACCCCAACCAAGTCAGGTAGTAAGTATGATGAGAGCCGTG 120
 Db 91 GCACAAATTATCAGGAACACCCCAACCAAGTCAGGTAGTAAGTATGATGAGAGCCGTG 32
 QY 121 TTTCATGTTTAATTAATT 137
 Db 31 TTTCATGTTTAATTAATT 15
 RESULT 8
 AW950480/c

```

LOCUS      AW950480                696 bp    mRNA    linear    EST 01-JUN-2000
DEFINITION EST362550 MAGC resequencences, MAGA Homo sapiens cDNA, mRNA sequence.
ACCESSION  AW950480
VERSION    AW950480.1  GI:8140134
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 696)
AUTHORS   Hegde, P., Qi, R., Abernathy, K., Dharap, S., Gaspard, R., Gay, C.,
            Holt, I.E., Saeed, A.I., Sharov, V., Lee, N.H., Yeatman, T.J. and
            Quackenbush, J.
TITLE      Assessment of gene expression patterns in a model of colon tumor
            metastasis using a 19,200 element cDNA microarray
JOURNAL    Unpublished (2000)
COMMENT    Contact: John Quackenbush
            The Institute for Genomic Research
            9712 Medical Center Dr., Rockville, MD 20850, USA
            Tel: 301 838 3528
            Fax: 301 838 0208
            Email: johnq@tigr.org
            Plate: 15
            Seq primer: Reverse.
FEATURES   Location/Qualifiers
            source          1..696
                        /organism="Homo sapiens"
                        /mol_type="mRNA"
                        /db_xref="taxon:9606"
                        /clone_lib="MAGE resequencences, MAGA"
                        /note="Vector: pBluescriptSKm"
ORIGIN
Query Match      100.0%; Score 137; DB 10; Length 696;
Best Local Similarity 100.0%; Pred. No. 3e-57;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAAATACCGAATGCAAGTGGTAGAAGCTGTGAAGCTGTAAGTCAAGTGT 60
DB 572 ATTATAAGGAAAAAGAAAATACCGAATGCAAGTGGTAGAAGCTGTGAAGCTGTAAGTCAAGTGT 513

QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAAGTGAAGTGAAGTGAAGTGAAGT 120
DB 512 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAAGTGAAGTGAAGTGAAGTGAAGT 453

QY 121 TTTGATGTTAATAATT 137
DB 452 TTTGATGTTAATAATT 436

RESULT 9
AV725574/c
LOCUS      AV725574 HTC Homo sapiens cDNA clone HTCBD10 5', mRNA sequence.
DEFINITION AV725574
ACCESSION  AV725574
VERSION    AV725574.1  GI:10831099
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 710)
AUTHORS   Gu, Y., Peng, Y., Song, H., Huang, Q., Yang, Y., Gao, G., Xiao, H., Xu, X.,
            Li, N., Qian, B., Liu, F., Qu, J., Gao, X., Cheng, Z., Xu, Z., Zeng, L.,
            Xu, S., Gu, W., Tu, Y., Jia, J., Fu, G., Ren, S., Zhong, M., Lu, G., Hu, R.,
            Chen, J., Chen, Z., and Han, Z.
            Homo sapiens cDNA HTC Clones
            Unpublished (2000)
            Contact: Zeguang Han
            Chinese National Human Genome Center at Shanghai
            351 Guo Shoujing Road, Zhangjiang Hi-Tech Park, Pudong, Shanghai
            201203, P. R. China
            Tel: 86-21-50801919(ex.45)

```

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FEATURES   Location/Qualifiers
            source          1..710
                        /organism="Homo sapiens"
                        /mol_type="mRNA"
                        /db_xref="taxon:9606"
                        /clone="HTCBD10"
                        /tissue_type="Hypothalamus"
                        /dev_stage="Adult"
                        /lab_host="SOLR"
                        /clone_lib="HTC"
                        /notes="Vector: pBluescript sk(-); Site_1: EcoRI; Site_2:
                        XhoI"
ORIGIN
Query Match      100.0%; Score 137; DB 9; Length 710;
Best Local Similarity 100.0%; Pred. No. 3e-57;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAAGAAAATACCGAATGCAAGTGGTAGAAGCTGTGAAGCTGTAAGTCAAGTGT 60
DB 396 ATTATAAGGAAAAAGAAAATACCGAATGCAAGTGGTAGAAGCTGTGAAGCTGTAAGTCAAGTGT 337

QY 61 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAAGTGAAGTGAAGTGAAGTGAAGT 120
DB 336 GCACAAATTATCAGGAACACCCCAAAACCAAGTGGTAGAAGTGAAGTGAAGTGAAGTGAAGT 277

QY 121 TTTGATGTTAATAATT 137
DB 276 TTTGATGTTAATAATT 260

RESULT 10
BG567176/c
LOCUS      BG567176 Homo sapiens cDNA clone IMAGE:4723833 5',
DEFINITION BG567176 Homo sapiens cDNA clone IMAGE:4723833 5',
            mRNA sequence.
ACCESSION  BG567176
VERSION    BG567176.1  GI:13574829
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 727)
AUTHORS   NIH-MGC http://mgi.nci.nih.gov/.
            National Institutes of Health, Mammalian Gene Collection (MGC)
            Unpublished (1999)
            Contact: Robert Strausberg, Ph.D.
            Email: cgapbs@mail.nih.gov
            Tissue Procurement: CLONETECH Laboratories, Inc.
            cDNA Library Preparation: CLONETECH Laboratories, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            http://image.llnl.gov
            Plate: LICM1583 row: p column: 10
            High quality sequence stop: 694.
FEATURES   Location/Qualifiers
            source          1..727
                        /organism="Homo sapiens"
                        /mol_type="mRNA"
                        /db_xref="taxon:9606"
                        /clone="IMAGE:4723833"
                        /lab_host="NIH MGC 76"
                        /notes="Organ: liver; Vector: pDNR-LIB (Clontech); Site 1:
                        SfiI (ggccctcgcc); Site 2: SfiI (ggccattggcc); 5' and
                        3' adaptors were used in cloning as follows: 5' adaptor
                        sequence: 5'-CACGCCCATATGCCC-3' and 3' adaptor sequence:

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Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAGGAAAGAAATAACCAATGACCAAGTGGTGAAGCTGGAAGTCAAGTGT 60
 DB 260 ATTATAGGAAAGAAATAACCAATGACCAAGTGGTGAAGCTGGAAGTCAAGTGT 201

QY 61 GCACAATTATCAGGAACACCCCAACCAACCAAGTGGTGAAGTCAAGTGTAGAGCCGCTG 120
 DB 200 GCACAATTATCAGGAACACCCCAACCAACCAAGTGGTGAAGTCAAGTGTAGAGCCGCTG 141

QY 121 TTGTGATGTTAATTAATT 137
 DB 140 TTGTGATGTTAATTAATT 124

RESULT 13
 BM546012/c
 LOCUS 1121 bp mRNA linear EST 20-FEB-2002
 DEFINITION AGENCOURT_6505286 NIH_MGC_125 Homo sapiens cDNA clone IMAGE:5588419
 5', mRNA sequence.
 ACCESSION BM546012
 VERSION BM546012
 KEYWORDS EST.
 SOURCE GI:18778623
 ORGANISM Homo sapiens (human)
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 1121)
 AUTHORS NIH-MGC http://mgi.nci.nih.gov/
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 Tissue Procurement: Invitrogen
 cDNA Library Preparation: Life Technologies, Inc.
 CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov
 Plate: LMAW12359 row: h column: 20
 High quality sequence start: 50
 High quality sequence stop: 764.
 FEATURES
 Location/Qualifiers
 1..1121
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:5588419"
 /lab_host="DH10B"
 /clone_lib="NIH MGC 125"
 /note="Organ: ovary (pool of 3); Vector: pCMV-SPORT6;
 Site 1: EcoRV (destroyed); Site 2: NotI; RNA source pool
 of three ovaries, from females ranging in age from 38 to
 49 yo. Library is oligo-dr primed and directionally cloned
 (EcoRV site is destroyed upon cloning). Average insert
 size 2.1 kb, insert size range 1-3.5 kb. Library is
 normalized and enriched for full-length clones and was
 constructed by C. Gruber (Invitrogen). Research Genetics
 tracking code 036."

ORIGIN
 Query Match 100.0%; Score 137; DB 12; Length 1121;
 Best Local Similarity 100.0%; Pred. No. 3.1e-57;
 Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAGGAAAGAAATAACCAATGGAAGTGGTGAAGCTGGAAGTCAAGTGT 60
 DB 514 ATTATAGGAAAGAAATAACCAATGGAAGTGGTGAAGCTGGAAGTCAAGTGT 455

QY 61 GCACAATTATCAGGAACACCCCAACCAACCAAGTGGTGAAGTCAAGTGTAGAGCCGCTG 120
 DB 454 GCACAATTATCAGGAACACCCCAACCAACCAAGTGGTGAAGTCAAGTGTAGAGCCGCTG 395

QY 121 TTGTGATGTTAATTAATT 137
 DB 394 TTGTGATGTTAATTAATT 378

RESULT 14
 CD110642/c
 LOCUS 743 bp mRNA linear EST 15-MAY-2003
 DEFINITION AGENCOURT_13995397 NIH_MGC_187 Homo sapiens cDNA clone
 IMAGE:30373580 5', mRNA sequence.
 ACCESSION CD110642
 VERSION CD110642.1
 KEYWORDS EST.
 SOURCE GI:30754851
 ORGANISM Homo sapiens (human)
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 743)
 AUTHORS NIH-MGC http://mgi.nci.nih.gov/
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgapbs-r@mail.nih.gov
 Tissue Procurement: Dr. Michael Brownstein and Dr. Miklos Palkovits
 CDNA Library Preparation: CLONTECH Laboratories, Inc.
 CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 http://image.llnl.gov
 Plate: NDCM167 row: j column: 21
 High quality sequence stop: 578.
 FEATURES
 Location/Qualifiers
 1..743
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:30373580"
 /lab_host="DH10B (T1 phage-resistant)"
 /clone_lib="NIH MGC 187"
 /note="Organ: Blood vessels - aorta, basilar and artery;
 Vector: pDNR-LIB; Site 1: SfiI (ggccattggcc); Site 2:
 SfiI (ggcgctcgccg); 5' and 3' adaptors were used in
 cloning as follows: 5' adaptor sequence:
 5'-CACGACCAATTATGCGC-3' and 3' adaptor sequence:
 5'-ATTCTAGAGCCGCGCGCCGCGCCGACATG-DT(30)BN-3' (where B = A,
 C, or G and N = A, C, G, or T). Average insert size 1.4 kb
 (range 0.5-4.0 kb). 14/15 colonies contained inserts by
 PCR. This library was enriched for full-length clones and
 was constructed by Clontech Laboratories (Palo Alto, CA).
 Note: this is a NIH_MGC Library."

ORIGIN
 Query Match 99.3%; Score 136; DB 14; Length 743;
 Best Local Similarity 100.0%; Pred. No. 9.5e-57;
 Matches 136; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAGGAAAGAAATAACCAATGGAAGTGGTGAAGCTGGAAGTCAAGTGT 60
 DB 138 ATTATAGGAAAGAAATAACCAATGGAAGTGGTGAAGCTGGAAGTCAAGTGT 79

QY 61 GCACAATTATCAGGAACACCCCAACCAACCAAGTGGTGAAGTCAAGTGTAGAGCCGCTG 120
 DB 78 GCACAATTATCAGGAACACCCCAACCAACCAAGTGGTGAAGTCAAGTGTAGAGCCGCTG 19

QY 121 TTGTGATGTTAATTAATT 136
 DB 18 TTGTGATGTTAATTAATT 3

RESULT 15
 AW385154

LOCUS AW385154 352 bp mRNA linear EST 04-FEB-2000
 DEFINITION PM2-HT0451-281299-001-a07 HT0451 Homo sapiens cDNA, mRNA sequence.
 ACCESSION AW385154
 VERSION AW385154.1 GI:6889813
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 352)
 HCGP <http://www.ludwig.org.br/ORESTES>.
 TITLE The FAPESP/LICR Human Cancer Genome Project
 JOURNAL Unpublished (1999)
 COMMENT Contact: Simpson A.J.G.
 Laboratory of Cancer Genetics
 Ludwig Institute for Cancer Research
 Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
 Brazil
 Tel: +55-11-2704922
 Fax: +55-11-2707001
 Email: asimpson@ludwig.org.br
 This sequence was derived from the FAPESP/LICR Human Cancer Genome
 Project. This entry can be seen in the following URL
 (<http://www.ludwig.org.br/scripts/gethtml2.pl?ti=PM2&t2=PM2-HT0451-281299-001-a07&t3=1999-12-28&t4=1>)
 Seq primer: puc 18 forward
 High quality sequence stop: 352.

FEATURES

1..352
 Location/Qualifiers
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /dev_stage="Adult"
 /clone_lib="HT0451"
 /note="Organ: head neck; Vector: puc18; Site 1: Sma1;
 Site 2: Sma1; A mini-library was made by cloning products
 derived from ORESTES PCR (U.S. Letters Patent application
 No. 196,716 - Ludwig Institute for Cancer Research)
 profiles into the pUC 18 vector. Reverse transcription of
 tissue mRNA and cDNA amplification were performed under
 low stringency conditions."

ORIGIN

Query Match 87.6%; Score 120; DB 10; Length 352;
 Best Local Similarity 100.0%; Pred. No. 7.9e-49;
 Matches 120; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 18 AATAACGCAATGCAAGTGGTGAAGCTGTGAACCTCAGGTGTGCACAAATTATCAGGAAC 77
 Db 1 AATAACGCAATGCAAGTGGTGAAGCTGTGAACCTCAGGTGTGCACAAATTATCAGGAAC 60
 QY 78 ACCCCAAACCAAAGTCAGTGAATAGCATGAGAGCCGCTTTGATGTTAATTAATT 137
 Db 61 ACCCCAAACCAAAGTCAGTGAATAGCATGAGAGCCGCTTTGATGTTAATTAATT 120

Search completed: April 6, 2004, 17:40:07
 Job time : 506.195 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 13:02:17 ; Search time 574.481 Seconds
(without alignments)
10336.278 Million cell updates/sec

Title: US-09-966-264D-1
Perfect score: 137
Sequence: 1 attataaggaagaaagaaaa.....gtgtttgatgttaattaatt 137

Scoring table: OLIGO NUC
Gapop 60.0 , Gapext 60.0

Searched: 3470272 seqs, 21671516995 residues

Word size : 0

Total number of hits satisfying chosen parameters: 6940544

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : GenEmbl.*

- 1: gb_ba.*
- 2: gb_hgt.*
- 3: gb_in.*
- 4: gb_cm.*
- 5: gb_ov.*
- 6: gb_pat.*
- 7: gb_ph.*
- 8: gb_pl.*
- 9: gb_pr.*
- 10: gb_ro.*
- 11: gb_sts.*
- 12: gb_sy.*
- 13: gb_un.*
- 14: gb_vi.*
- 15: em_ba.*
- 16: em_fun.*
- 17: em_hum.*
- 18: em_in.*
- 19: em_mu.*
- 20: em_om.*
- 21: em_or.*
- 22: em_ov.*
- 23: em_pat.*
- 24: em_ph.*
- 25: em_pl.*
- 26: em_ro.*
- 27: em_sts.*
- 28: em_un.*
- 29: em_vi.*
- 30: em_hgt_hum.*
- 31: em_hgt_inv.*
- 32: em_hgt_other.*
- 33: em_hgt_mus.*
- 34: em_hgt_pln.*
- 35: em_hgt_rod.*
- 36: em_hgt_man.*
- 37: em_hgt_vrt.*
- 38: em_sy.*
- 39: em_hgtgo_hum.*
- 40: em_hgtgo_mus.*
- 41: em_hgtgo_other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
C 1	137	100.0	2148	11	G15848	G15848 human STS C
C 2	137	100.0	2563	9	AK129855	AK129855 Homo sapi
C 3	137	100.0	2691	6	AX538618	AX538618 Sequence
C 4	137	100.0	4658	9	BC028720	BC028720 Homo sapi
C 5	137	100.0	8689	6	AX538622	AX538622 Sequence
C 6	137	100.0	11443	6	AX538634	AX538634 Sequence
C 7	137	100.0	13957	6	AX409637	AX409637 Sequence
C 8	137	100.0	13957	6	AX538581	AX538581 Sequence
C 9	137	100.0	13957	9	HUMDYS	M18533 Homo sapien
C 10	137	100.0	98056	9	AC006061	AC006061 Homo sapi
C 11	117	85.4	13977	6	AR220819	AR220819 Sequence
C 12	65	47.4	912120	2	AC023414	AC023414 Homo sapi
C 13	46	33.6	13887	4	AF070485	AF070485 Canis fam
C 14	46	33.6	272578	2	AC108338	AC108338 Rattus no
C 15	46	33.6	272578	2	AC114184	AC114184 Rattus no
C 16	42	30.7	13815	6	AX306153	AX306153 Sequence
C 17	42	30.7	13815	6	AX538582	AX538582 Sequence
C 18	42	30.7	13815	10	MUSDYSA	M68859 Mouse cysr
C 19	42	30.7	19307	6	AR093392	AR093392 Sequence
C 20	42	30.7	19307	6	AR142592	AR142592 Sequence
C 21	42	30.7	189131	10	AL645477	AL645477 Mouse DNA
C 22	30	21.9	13575	5	GGDYS	X13369 Chicken mRN
C 23	22	16.1	208367	2	EX682542	EX682542 Mus muscu
C 24	21	15.3	157446	2	AC118861	AC118861 Rattus no
C 25	20	14.6	99871	9	AL603824	AL603824 Human DNA
C 26	20	14.6	94605	9	AF410480S2	AF360695 Homo sapi
C 27	20	14.6	114226	2	AC094710	AC094710 Plasmodiu
C 28	20	14.6	136467	9	AF487553	AF487553 Homo sapi
C 29	20	14.6	143410	9	AC087636	AC087636 Homo sapi
C 30	20	14.6	145990	10	AC117837	AC117837 Mus muscu
C 31	20	14.6	157216	2	AC008244	AC008244 Homo sapi
C 32	20	14.6	159557	9	AC127898	AC127898 Homo sapi
C 33	20	14.6	160750	9	HSJ177110	AL080284 Human DNA
C 34	20	14.6	161601	2	AC021147	AC021147 Homo sapi
C 35	20	14.6	162612	2	AC118722	AC118722 Mus muscu
C 36	20	14.6	165660	2	AC027194	AC027194 Homo sapi
C 37	20	14.6	165789	2	AL591029	AL591029 Homo sapi
C 38	20	14.6	166782	2	AC021516	AC021516 Homo sapi
C 39	20	14.6	169524	2	AC023111	AC023111 Homo sapi
C 40	20	14.6	174398	9	AC009988	AC009988 Homo sapi
C 41	20	14.6	177185	9	AC016632	AC016632 Homo sapi
C 42	20	14.6	180702	2	AC121392	AC121392 Rattus no
C 43	20	14.6	181390	9	AC074237	AC074237 Homo sapi
C 44	20	14.6	182126	2	AC016639	AC016639 Homo sapi
C 45	20	14.6	187466	9	AC016322	AC016322 Homo sapi

ALIGNMENTS

RESULT 1
G15848/c
LOCUS
DEFINITION
human STS CHLC UTR_01924_M18533. P56108 clone UTR_01924_M18533,
linear STS 19-JAN-1996
sequence tagged site.
ACCESSION
G15848.1
VERSION
G15848.1
KEYWORDS
STS; STS sequence; primer; sequence tagged site.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 2148)
Murray,J., Sheffield,V, Weber,J.L., Duyk,G. and Buetow,K.H.
Cooperative Human Linkage Center

JOURNAL
COMMENT
Unpublished (1995)
Synonyms: UTR_01924_M18533, CHLC.UTR_01924_M18533.T36152
Contact: Dr. Jeffrey C. Murray

UofI
The University of Iowa
Department of Pediatrics, Iowa City, IA 52242, USA
Tel: (319) 356-3508
Fax: (319) 356-3347
Email: jeff-murray@uiowa.edu

Primer A: AACGATTTTGGGTGTTTA
Primer B: GATATCAGCCCAAGGATG
STS size: 189
PCR Profile:

denature: 30 seconds at 94 degrees C
annealing: 75 seconds at 55 degrees C
extension: 15 seconds at 72 degrees C
PCR cycles: 27
extension: 6 minutes at 72 degrees C

Protocol:
Template: 30mg genomic DNA
Primer: each 1.5 pmole
dNTPs: each 200 uM
Taq Polymerase: 0.3 units
Total Vol: 10 ul

Buffer:
MgCl2: 1.5mM
KCl: 50mM
Tris: 10mM
pH: 8.3

Prepared with primer pairs derived from M18533.

FEATURES
source
Location/Qualifiers
1..2148
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
350..538
primer_bind 350..369
primer_bind complement(519..538)

Query Match 100.0%; Score 137; DB 11; Length 2148;
Best Local Similarity 100.0%; Pred. No. 5.4e-63;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTATAAGGAAAAGAAAATACGCAATCGCAAGTGGTGAAGCTGTGAAGCTCAGGTGT 60
Db 2033 ATTATAAGGAAAAGAAAATACGCAATCGCAAGTGGTGAAGCTGTGAAGCTCAGGTGT 1974
Qy 61 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAGGTAGATAATAGCAGAGCCGCGT 120
Db 1973 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAGGTAGATAATAGCAGAGCCGCGT 1914
Qy 121 TTTGATGTTAATTAATT 137
Db 1913 TTTGATGTTAATTAATT 1897

RESULT 2
AK129855/c AK129855 2563 bp mRNA linear PRI 10-SEP-2003
LOCUS Homo sapiens cDNA FLJ26345 fis, clone HRT03668.

DEFINITION AK129855
ACCESSION AK129855
VERSION AK129855.1 GI:34526478
KEYWORDS oligo capping; fis (full insert sequence).

SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1
Ota,T., Nakagawa,S., Seroh,A., Mizuguchi,H., Inagaki,H., Suzuki,Y.,

Hata,H., Nakagawa,K., Mizuno,S., Morinaga,M., Kawamura,M.,
Sugiyama,T., Irie,R., Otsuki,T., Sato,H., Nishikawa,T.,
Sugiyama,A., Kawakami,B., Nagai,K., Isogai,T. and Sugano,S.
NEDO human cDNA sequencing project

TITLE Unpublished
JOURNAL 2 (bases 1 to 2563)
REFERENCE Sugano,S. and Suzuki,Y.
AUTHORS Direct Submission
TITLE Submitted (31-JUL-2003) Sumio Sugano, Institute of Medical Science,
JOURNAL University of Tokyo, Laboratory of Genome Structure, Human Genome
Center, Shirokane-dai, 4-6-1, Minato-ku, Tokyo 108-8639, Japan
(E-mail: filcdna@u-tokyo.ac.jp, Tel:81-3-5449-5286,
Fax:81-3-5449-5416)

COMMENT NEDO human cDNA sequencing project supported by Ministry of
Economy, Trade and Industry of Japan; cDNA full insert sequencing:
Research Association for Biotechnology (RAB); cDNA library
construction and 5'-end one pass sequencing: Institute of Medical
Science, University of Tokyo, Laboratory of Genome Structure, Human
Genome Center; 3'-end one pass sequencing: RAB; clone selection for
full insert sequencing: RAB and Helix Research Institute.

FEATURES
source
Location/Qualifiers
1..2563
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="HRT03668"
/tissue_type="heart"
/clone_lib="HRT"
/note="cloning vector: pME18SFL3"

ORIGIN
Query Match 100.0%; Score 137; DB 9; Length 2563;
Best Local Similarity 100.0%; Pred. No. 5.4e-63;
Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATTATAAGGAAAAGAAAATACGCAATCGCAAGTGGTGAAGCTGTGAAGCTCAGGTGT 60
Db 1905 ATTATAAGGAAAAGAAAATACGCAATCGCAAGTGGTGAAGCTGTGAAGCTCAGGTGT 1846
Qy 61 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAGGTAGATAATAGCAGAGCCGCGT 120
Db 1845 GCACAATTATCAGGAACACCCCAAAACCAAAAGTGAGGTAGATAATAGCAGAGCCGCGT 1786
Qy 121 TTTGATGTTAATTAATT 137
Db 1785 TTTGATGTTAATTAATT 1769

RESULT 3
AX538618/c AX538618 2591 bp DNA linear PAT 23-NOV-2002
LOCUS Sequence 38 from Patent WO0229056.
DEFINITION AX538618
ACCESSION AX538618
VERSION AX538618.1 GI:25271161
KEYWORDS
SOURCE Homo sapiens (human)

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1
AUTHORS Chamberlain,J.S. and Harper,S.Q.
TITLE Mini-dystrophin nucleic acid and peptide sequences
JOURNAL Patent, WO 0229056-A 38 11-APR-2002.
THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)

FEATURES
source
Location/Qualifiers
1..2691
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"

ORIGIN
Query Match 100.0%; Score 137; DB 6; Length 2691;
Best Local Similarity 100.0%; Pred. No. 5.4e-63;

artificial sequences.					
REFERENCE	1				
AUTHORS	Chamberlain,J.S. and Harper,S.Q.				
TITLE	Mini-dystrophin nucleic acid and peptide sequences				
JOURNAL	Patent: WO 0229056-A 42 11-APR-2002;				
KEYWORDS	THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)				
FEATURES	Location/Qualifiers				
source	1..8689				
	/organism="synthetic construct"				
	/mol_type="unassigned DNA"				
	/db_xref="taxon:32630"				
	/note="Synthetic"				
ORIGIN					
Query Match	100.0%; Score 137; DB 6; Length 8689;				
Best Local Similarity	100.0%; Pred. No. 5.1e-63;				
Matches 137; Conservative	0; Mismatches 0; Indels 0; Gaps 0;				
QY	1 ATTATAAGGAAAAAGAAATTAACCAATGCACAAGTGGTGGAAGCTGTGAAGCTGTGAAGCTCAGGTGT 60				
DB	8031 ATTATAAGGAAAAAGAAATTAACCAATGCACAAGTGGTGGAAGCTGTGAAGCTGTGAAGCTCAGGTGT 7972				
QY	61 GCACAATTATCAGGAACACCCCAAAACCACCAAGTCAGGTAGAAAATAGCATGAGAAGCCCGTG 120				
DB	7971 GCACAATTATCAGGAACACCCCAAAACCACCAAGTCAGGTAGAAAATAGCATGAGAAGCCCGTG 7912				
QY	121 TTTCATGTTTAATTAATT 137				
DB	7911 TTTCATGTTTAATTAATT 7895				
RESULT 6					
AX538624/C					
LOCUS	AX538624 11443 bp DNA linear PAT 23-NOV-2002				
DEFINITION	Sequence 44 from Patent WO0229056.				
ACCESSION	AX538624				
VERSION	AX538624.1 GI:25271175				
KEYWORDS	synthetic construct				
SOURCE	synthetic construct				
ORGANISM	artificial sequences.				
REFERENCE	1				
AUTHORS	Chamberlain,J.S. and Harper,S.Q.				
TITLE	Mini-dystrophin nucleic acid and peptide sequences				
JOURNAL	Patent: WO 0229056-A 44 11-APR-2002;				
KEYWORDS	THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)				
FEATURES	Location/Qualifiers				
source	1..11443				
	/organism="synthetic construct"				
	/mol_type="unassigned DNA"				
	/db_xref="taxon:32630"				
	/note="Synthetic"				
ORIGIN					
Query Match	100.0%; Score 137; DB 6; Length 11443;				
Best Local Similarity	100.0%; Pred. No. 5e-63;				
Matches 137; Conservative	0; Mismatches 0; Indels 0; Gaps 0;				
QY	1 ATTATAAGGAAAAAGAAATTAACCAATGCACAAGTGGTGGAAGCTGTGAAGCTGTGAAGCTCAGGTGT 60				
DB	10785 ATTATAAGGAAAAAGAAATTAACCAATGCACAAGTGGTGGAAGCTGTGAAGCTGTGAAGCTCAGGTGT 10726				
QY	61 GCACAATTATCAGGAACACCCCAAAACCACCAAGTCAGGTAGAAAATAGCATGAGAAGCCCGTG 120				
DB	10725 GCACAATTATCAGGAACACCCCAAAACCACCAAGTCAGGTAGAAAATAGCATGAGAAGCCCGTG 10666				
QY	121 TTTCATGTTTAATTAATT 137				
DB	10665 TTTCATGTTTAATTAATT 10649				
RESULT 7					
AX409637/G					
LOCUS	AX409637 13957 bp DNA linear PAT 14-JUN-2002				
DEFINITION	Sequence 2284 from Patent WO0229103.				
ACCESSION	AX409637				
VERSION	AX409637.1 GI:21442342				
KEYWORDS	Homo sapiens (human)				
SOURCE	Homo sapiens				
ORGANISM	Homo sapiens				
REFERENCE	1				
AUTHORS	Alvares,C., Horne,D., Peres-da-Silva,S. and Vockley,J.G.				
TITLE	Gene expression profiles in liver cancer				
JOURNAL	Patent: WO 0229103-A 2284 11-APR-2002;				
KEYWORDS	GENE LOGIC INC (US)				
FEATURES	Location/Qualifiers				
source	1..13957				
	/organism="Homo sapiens"				
	/mol_type="unassigned DNA"				
	/db_xref="taxon:9606"				
	/notes="EMBL/GenBank Accession No. M18533"				
ORIGIN					
Query Match	100.0%; Score 137; DB 6; Length 13957;				
Best Local Similarity	100.0%; Pred. No. 5e-63;				
Matches 137; Conservative	0; Mismatches 0; Indels 0; Gaps 0;				
QY	1 ATTATAAGGAAAAAGAAATTAACGCAATGGCAAGTGGTGGAAGCTGTGAAGCTGTGAAGCTCAGGTGT 60				
DB	13299 ATTATAAGGAAAAAGAAATTAACGCAATGGCAAGTGGTGGAAGCTGTGAAGCTGTGAAGCTCAGGTGT 13240				
QY	61 GCACAATTATCAGGAACACCCCAAAACCACCAAGTCAGGTAGAAAATAGCATGAGAAGCCCGTG 120				
DB	13239 GCACAATTATCAGGAACACCCCAAAACCACCAAGTCAGGTAGAAAATAGCATGAGAAGCCCGTG 13180				
QY	121 TTTCATGTTTAATTAATT 137				
DB	13179 TTTCATGTTTAATTAATT 13163				
RESULT 8					
AX538581/C					
LOCUS	AX538581 13957 bp DNA linear PAT 23-NOV-2002				
DEFINITION	Sequence 1 from Patent WO0229056.				
ACCESSION	AX538581				
VERSION	AX538581.1 GI:25271086				
KEYWORDS	Homo sapiens (human)				
SOURCE	Homo sapiens				
ORGANISM	Homo sapiens				
REFERENCE	1				
AUTHORS	Chamberlain,J.S. and Harper,S.Q.				
TITLE	Mini-dystrophin nucleic acid and peptide sequences				
JOURNAL	Patent: WO 0229056-A 1 11-APR-2002;				
KEYWORDS	THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)				
FEATURES	Location/Qualifiers				
source	1..13957				
	/organism="Homo sapiens"				
	/mol_type="unassigned DNA"				
	/db_xref="taxon:9606"				
ORIGIN					
Query Match	100.0%; Score 137; DB 6; Length 13957;				
Best Local Similarity	100.0%; Pred. No. 5e-63;				
Matches 137; Conservative	0; Mismatches 0; Indels 0; Gaps 0;				
QY	1 ATTATAAGGAAAAAGAAATTAACGCAATGGCAAGTGGTGGAAGCTGTGAAGCTGTGAAGCTCAGGTGT 60				
DB	13299 ATTATAAGGAAAAAGAAATTAACGCAATGGCAAGTGGTGGAAGCTGTGAAGCTGTGAAGCTCAGGTGT 13240				
QY					

RESULT 10

AC006061 Homo sapiens X BAC GSHB-19024 (Genome Systems Human BAC Library) linear PRI 01-MAY-2002
 LOCUS AC006061 complete sequence.
 DEFINITION AC006061
 ACCESSION AC006061.1 GI:4204246
 VERSION
 KEYWORDS
 SOURCE Homo sapiens (human)
 ORGANISM
 Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 98056)
 Muzny, D., Aronson, A.D., Bouck, J., Brundage, E., Bunac, C., Chen, Z.,
 Di, W., Ding, Y., Dugan, S., Durbin, J., Forcum, J., Garcia, C.,
 Gorrell, J.H., Gorrell, L.L., Hernandez, J., Jackson, L.,
 Kondejewski, N., Leal, B., Lichtarge, O., Liu, W., Logan, O., Lu, J.,
 Martinez, C., Oswal, G., Pampall, L.R., Parish, B.J., Perez, L.,
 Rasid, N.D., Rives, C., Scherer, S.E., Shen, H., Simon, M., Vo, Q.,
 Williamson, A., Worley, K.C., Yu, W., Zhou, X., Nelson, D., and
 Gibbs, R.A.
 Direct Submission
 Unpublished
 2 (bases 1 to 98056)
 Worley, K.C.
 Direct Submission
 Submitted (26-NOV-1998) Molecular and Human Genetics, Baylor
 College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 3 (bases 1 to 98056)
 Worley, K.C.
 Direct Submission
 Submitted (30-JAN-1999) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 4 (bases 1 to 98056)
 Worley, K.C.
 Direct Submission
 Submitted (02-FEB-1999) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 5 (bases 1 to 98056)
 Worley, K.C.
 Direct Submission
 Submitted (04-FEB-1999) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 6 (bases 1 to 98056)
 Worley, K.C.
 Direct Submission
 Submitted (28-MAR-2001) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 7 (bases 1 to 98056)
 Worley, K.C.
 Direct Submission
 Submitted (07-MAR-2002) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 8 (bases 1 to 98056)
 Worley, K.C.
 Direct Submission
 Submitted (01-MAY-2002) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA
 On Jan 30, 1999 this sequence version replaced gi:4176317.
 INFORMATION: <http://gc.bcm.tmc.edu:8088/home.html> or email
 gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the
 entire insert of this clone. Overlapping regions of clones are only
 sequenced and submitted once, so the sequence for the remainder of
 the insert may be found in the record for the adjacent clones.

Overlapping clones are noted at the beginning and end of the
 Features listing.

ANNOTATION OF FEATURES:

STSs are identified using ePCR (Genome Res. 7:541-550) searches
 of a local database that includes entries from dbSTS, GDB, and
 local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green,
 unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST
 (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the
 EST and cDNA sequences. Genes demonstrate at least two exons
 flanked by consensus splice sites that maintained sequence
 continuity across the splice junctions. Sequences that are not
 identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum
 standard of double strand coverage with a minimum of 2 clones and 2
 reads with no ambiguities or 2 chemistries with a minimum of 2
 clones and 3 reads with no ambiguities. If the sequence quality for
 a region does not meet this standard, it will be indicated in the
 annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality
 standards - estimated error rate less than 1 per 10,000 bases.

FEATURES

Source	Location/Qualifiers
1..98056	/organism="Homo sapiens"
	/mol_type="genomic DNA"
	/db_xref="taxon:9606"
	/chromosome="X"
	/clone="GSHB-19024"
1..1999	/notes="overlaps bases 151178..153177 of clone AC117405"
	/function="clone overlap"
repeat_region	40..82
	/rpt_family="L2"
repeat_region	378..507
	/rpt_family="L1MD3"
repeat_region	complement(518..570)
	/rpt_family="(GGA)n"
repeat_region	571..729
	/rpt_family="L1MC4"
repeat_region	730..987
	/rpt_family="MSTD"
repeat_region	complement(991..1400)
	/rpt_family="L2"
repeat_region	complement(1649..1779)
	/rpt_family="MIR"
repeat_region	complement(1783..2212)
	/rpt_family="LTR16A"
repeat_region	complement(3626..3656)
	/rpt_family="(GA)n"
repeat_region	complement(4713..5074)
	/rpt_family="THE1B"
repeat_region	5702..7108
	/rpt_family="L2"
repeat_region	complement(8670..8734)
	/rpt_family="MLTIG"
repeat_region	complement(9069..9144)
	/rpt_family="MLTIG"
repeat_region	complement(9254..9351)
	/rpt_family="MIR"
repeat_region	complement(9537..9688)
	/rpt_family="MER5A"
repeat_region	9797..9942
	/rpt_family="MIR"
repeat_region	complement(10403..10429)
	/rpt_family="AT rich"
repeat_region	complement(10673..10705)
	/rpt_family="(CA)n"
repeat_region	complement(11460..11755)
	/rpt_family="Alu5x"


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repeat_region 11977..12125
/rpt_family="MER5A"
12556..11643
/rpt_family="L1MD2"
13618..13808
/rpt_family="L1MD3"
complement(11997..14297)
/rpt_family="MER4D"
14327..14625
/rpt_family="AluSg"
complement(14548..14915)
/rpt_family="MER4D"
complement(14915..15147)
/rpt_family="MER4D"
15286..15404
/rpt_family="FLAM_C"
16169..16463
/rpt_family="AluSx"
16520..16570
/rpt_family="L2"
16688..16904
/rpt_family="L2"
complement(16955..17147)
/rpt_family="MIR"
complement(17171..17467)
/rpt_family="AluY"
complement(20101..20241)
/rpt_family="MIR"
22625..22788
/standard_name="DXS1097"
/db_xref="GDB:190032"
24144..24435
/rpt_family="AluSg"
complement(25010..25235)
/rpt_family="L2"
25643..25771
/rpt_family="MER58A"
complement(26111..26429)
/rpt_family="L2"
complement(26578..26636)
/rpt_family="(CATA)n"
complement(26956..27027)
/rpt_family="L1MA9"
complement(27038..27188)
/rpt_family="L2"
27203..27641
/rpt_family="L1MA8"
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Matches 137; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATTATAAGGAAAAGAAAATAACGCAATGCGAAGTGTGAGTGTGAACTCAGGTGT 60
Db 91659 ATTATAAGGAAAAGAAAATAACGCAATGCGAAGTGTGAGTGTGAACTCAGGTGT 91718

QY 61 GCACATTATCAGGACACCCCAACCAAGTGTGAGTGTGAACTCAGGACCCG 120
Db 91719 GCACATTATCAGGACACCCCAACCAAGTGTGAGTGTGAACTCAGGACCCG 91778

QY 121 TTGTGATGTTAATTAATT 137
Db 91779 TTGTGATGTTAATTAATT 91795

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RESULT 11
AR220819/c 13977 bp DNA linear PAT 26-SEP-2002
LOCUS Sequence 60 from patent US 6426186.
DEFINITION AR220819
ACCESSION AR220819
VERSION AR220819.1 GI:23327696
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
Unclassified.
1 (bases 1 to 13977)
AUTHORS Jones,K.A., Volkmut,W. and Walker,M.G.
TITLE Bone remodeling genes
JOURNAL Patent: US 6426186-A 60 30-JUL-2002;
FEATURES
Location/Qualifiers
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ORIGIN
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QY 61 GCACATTATCAGGACACCCCAACCAAGTGTGAGTGTGAACTCAGGACCC 117
Db 13246 GCACATTATCAGGACACCCCAACCAAGTGTGAGTGTGAACTCAGGACCC 13190

RESULT 12
AC023414 212120 bp DNA linear HTG 11-APR-2000
LOCUS Homo sapiens chromosome 14 clone RP11-767B5 map 14, WORKING DRAFT
DEFINITION SEQUENCE, 31 unordered pieces.
ACCESSION AC023414.2 GI:7534009
VERSION HTG; HTGS_PHASE1; HTGS_DRAFT.
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 212120)
AUTHORS Birren,B., Linton,L., Nusbaum,C. and Lander,E.
TITLE Homo sapiens chromosome 14, clone RP11-767B5
JOURNAL Unpublished
REFERENCE
2 (bases 1 to 212120)
AUTHORS Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
Anderson,S., Baldwin,J., Barna,N., Beda,F., Boguslavsky,L.,
Bouckgalter,B., Brown,A., Burkett,G., Campopiano,A., Castle,A.,
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Grand-Pierre,N., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A.,
Klein,J., Landers,T., Largocque,K., Lehoczy,J., Levine,R.,
Lieu,C., Liu,G., Locke,K., Macdonald,P., Marquis,N., McCarthy,M.,
McEwan,P., McGurk,A., McKernan,K., McPheters,R., Meldrum,J.,
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Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Olivari,T.M.,
Peterson,K., Pierre,N., Pisani,C., Pollara,V., Raymond,C.,
Riley,R., Rogov,P., Rothman,D., Roy,A., Santos,R., Schauer,S.,
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Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J., Zimmer,A. and
Zody,M.

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Direct Submission
Submitted (14-FEB-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Apr 11, 2000 this sequence version replaced gi:6970579.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L6731
Center clone name: 767_B_5
----- Summary Statistics
Sequencing vector: M13, M77815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 185633 bases at least Q40
Consensus quality: 195115 bases at least Q30
Consensus quality: 201837 bases at least Q20
Insert size: 178000; agarose-fp
Insert size: 209120; sum-of-contents
Quality coverage: 5.8 in Q20 bases; agarose-fp
Quality coverage: 4.9 in Q20 bases; sum-of-contents

* NOTE: This is a 'working draft' sequence. It currently consists of 31 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

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*	2325	3345: contig of 1021 bp in length
*	3346	3445: gap of 100 bp
*	3446	4502: contig of 1057 bp in length
*	4503	5756: contig of 1154 bp in length
*	5757	5856: gap of 100 bp
*	5857	7097: contig of 1241 bp in length
*	7098	7197: gap of 100 bp
*	7198	8236: contig of 1039 bp in length
*	8237	8336: gap of 100 bp
*	8337	9353: contig of 1017 bp in length
*	9354	9453: gap of 100 bp
*	10705	contig of 1252 bp in length
*	10706	10805: gap of 100 bp
*	10806	12332: contig of 1527 bp in length
*	12333	12432: gap of 100 bp
*	12433	13877: contig of 1445 bp in length
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*	15202	15301: gap of 100 bp
*	15302	16968: contig of 1657 bp in length
*	16969	17068: gap of 100 bp
*	17069	19060: contig of 1592 bp in length
*	19061	19160: gap of 100 bp
*	19161	22639: contig of 3479 bp in length
*	22640	22739: gap of 100 bp
*	22740	26376: contig of 3637 bp in length
*	26377	26476: gap of 100 bp
*	26477	29275: contig of 2799 bp in length
*	29276	29375: gap of 100 bp
*	29376	33655: contig of 3890 bp in length
*	33656	33655: gap of 100 bp
*	33656	37012: contig of 3647 bp in length
*	37013	37112: gap of 100 bp

3390: contig of 2278 bp in length
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47621: contig of 3831 bp in length
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53852: contig of 6131 bp in length
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63422: contig of 9470 bp in length
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73361: contig of 13839 bp in length
7461: gap of 100 bp
92465: contig of 15004 bp in length
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106951: contig of 14386 bp in length
107051: gap of 100 bp
129678: contig of 22627 bp in length
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181939

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HTG, HTGS_PASEL1, HTGS_DRAFT; HTGS_ENRICHED.
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Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 279539)
Muzny D.Marie., Metzker M.Lee., Abramson S., Adams C., Alder J.,
Allen C., Allen H., Albrechts S., Amin A., Anguiano D.,
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Shetty J., Shwartsbeyn A., Sisson I., Sitter C.D., Smajls D.,
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Wang Q., Wang S., Warren J., Warren R., Wei X., White F.,
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Wright D., Wright R., Wu J., Yakub S., Yen J., Yoon L., Yoon V.,
Yu F., Zhang J., Zhou J., Zhou X., Zhao S., Dunn D., von
Niederhausern A., Weiss R., Smith D.R., Holt R.A., Smith H.O.,
Weinstock G. and Gibbs R.A.
Direct Submission
Unpublished
2 (bases 1 to 279539)
Worley K.C.
Direct Submission
Submitted (07-MAR-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 279539)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (11-OCT-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Oct 9, 2002 this sequence version replaced gi:21733949.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas

(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information

Center project name: GLPV
Center clone name: CH230-230F23
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 212054 bases at least Q40
Consensus quality: 219191 bases at least Q30
Consensus quality: 223518 bases at least Q20
Estimated insert size: 217504; sum-of-contigs estimation
Quality coverage: 5x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 21 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

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267565 267565: gap of unknown length
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271635 271635: contig of 3374 bp in length
275008 275008: gap of unknown length
275108 275108: gap of unknown length
277043 277043: contig of 1935 bp in length

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* 277044 277143: gap of unknown length
* 277144 278161: contig of 1018 bp in length
* 278162 278261: gap of unknown length
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FEATURES

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ORIGIN

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Query Match      33.6%; Score 46; DB 2; Length 279539;
Best Local Similarity 100.0%; Pred. No. 6.9e-14; Indels 0; Gaps 0;
Matches 46; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Search completed: April 6, 2004, 16:30:22
Job time : 581.481 secs

GenCore version 5.1.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 03:22:40 ; Search time 4175.64 Seconds
(without alignments)
10338.454 Million cell updates/sec

Title: US-09-966-264D-2

Perfect score: 996

Sequence: 1 ggggttgatgtagtagtaaa.....gtgttgatgtagtaataatt 996

Scoring table:

IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 3470272 seqs, 2167151695 residues

Total number of hits satisfying chosen parameters: 6940544

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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1: gb.ba.*

2: gb.htg.*

3: gb.in.*

4: gb.om.*

5: gb.ov.*

6: gb.pat.*

7: gb.ph.*

8: gb.pl.*

9: gb.pr.*

10: gb.ro.*

11: gb.sts.*

12: gb.sy.*

13: gb.un.*

14: gb.vi.*

15: em.ba.*

16: em.fun.*

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19: em.mu.*

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Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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c 2	858.2	86.2	212120	2	AC023414 Homo sapi
c 3	789.2	77.1	212120	2	AC023414 Homo sapi
c 4	435	43.7	189131	10	AL645477 Mouse DNA
5	431.4	43.3	272578	2	AC108338 Rattus no
6	412.2	41.4	2463	6	AX817316 Sequence
7	412.2	41.4	4658	9	BC028720 Homo sapi
8	410.8	41.2	5339	6	AX538620 Sequence
9	410.8	41.2	5417	6	AX538619 Sequence
10	410.8	41.2	5462	6	AX538621 Sequence
11	410.8	41.2	8689	6	AX538622 Sequence
12	410.8	41.2	11443	6	AX538624 Sequence
13	410.8	41.2	12057	6	AX538627 Sequence
14	410.8	41.2	12446	9	HSDMDR X14298 Human mRNA
15	410.8	41.2	13957	6	AX409637 Sequence
16	410.8	41.2	13957	6	AX538581 Sequence
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22	374.2	37.6	13815	6	AX306153 Sequence
23	374.2	37.6	13815	6	AX538582 Sequence
24	374.2	37.6	13815	10	MUSDYSA M68559 Mouse dysr
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28	350	35.1	350	9	AF213444 Homo sapi
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32	278.8	28.0	2563	9	AK129855 Homo sapi
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35	260	26.1	4075	6	E30221 Shortened d
36	260	26.1	4402	6	E30219 Shortened d
37	260	26.1	4402	6	E30220 Shortened d
38	252.8	25.4	3747	6	E30218 Shortened d
39	218	21.9	218	9	HUMDYS20 M6903 H. sapiens d
40	189.6	19.0	468	5	AF375546 Xenopus l
41	188.4	18.9	509	5	AF082741 Gallus ga
42	166.2	16.7	1534	5	AF375547 Scyliorhi
c 43	151.4	15.2	2148	11	G15848 G15848 human STS C
c 44	151.4	15.2	2563	9	AK129855 Homo sapi
c 45	151.4	15.2	2691	6	AX538618 Sequence

ALIGNMENTS

RESULT 1
AC006061/c AC006061 98056 bp DNA linear PRI 01-MAY-2002
LOCUS Homo sapiens X BAC GSHB-19024 (Genome Systems Human BAC Library)
DEFINITION complete sequence.
ACCESSION AC006061
VERSION HTG. AC006061.1 GI:4204246
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 98056)
AUTHORS Muzny,D., Aronson,A.D., Bouck,J., Brundage,E., Bunac,C., Chen,Z.,
Di,W., Ding,Y., Dugan,S., Durbin,J., Forcum,J., Garcia,C.,

Gorrell, J.H., Gorrell, L.L., Hernandez, J., Jackson, L.,
Kondejewski, N., Leal, B., Lichtarge, O., Liu, W., Logan, O., Lu, J.,
Martinez, C., Osval, G., Pampell, L.R., Parish, B.J., Perez, L.,
Rashid, N.D., Rives, C., Scherer, S.E., Shen, H., Simon, M., Vo, O.,
Williamson, A., Worley, K.C., Yu, W., Zhou, X., Nelson, D. and
Gibbs, R.A.
Direct Submission
Unpublished
2 (bases 1 to 98056)
Worley, K.C.
Direct Submission
Submitted (26-NOV-1998) Molecular and Human Genetics, Baylor
College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 98056)
Worley, K.C.
Direct Submission
Submitted (30-JAN-1999) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
4 (bases 1 to 98056)
Worley, K.C.
Direct Submission
Submitted (02-FEB-1999) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
5 (bases 1 to 98056)
Worley, K.C.
Direct Submission
Submitted (04-FEB-1999) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
6 (bases 1 to 98056)
Worley, K.C.
Direct Submission
Submitted (28-MAR-2001) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
7 (bases 1 to 98056)
Worley, K.C.
Direct Submission
Submitted (07-MAR-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
8 (bases 1 to 98056)
Worley, K.C.
Direct Submission
Submitted (01-MAY-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Jan 30, 1999 this sequence version replaced gi:4176317.
INFORMATION: <http://gc.bcm.tmc.edu:8088/home.html> or email
gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the
entire insert of this clone. Overlapping regions of clones are only
sequenced and submitted once, so the sequence for the remainder of
the insert may be found in the record for the adjacent clones.
Overlapping clones are noted at the beginning and end of the
Features listing.

ANNOTATION OF FEATURES:

STSs are identified using ePCR (Genome Res. 7:541-550) searches
of a local database that includes entries from dbSTS, GDB, and
local mapping efforts.
Repeats are identified using RepeatMasker (A. Smit and P. Green,
unpublished.) for Human and Mouse sequences.
Genes and Region of sequence similarity are identified by BLAST
(Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the
EST and cDNA sequences. Genes demonstrate at least two exons
flanked by consensus splice sites that maintained sequence
continuity across the splice junctions. Sequences that are not
identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum
standard of double strand coverage with a minimum of 2 clones and 2
reads with no ambiguities or 2 chemistries with a minimum of 2
clones and 3 reads with no ambiguities. If the sequence quality for
a region does not meet this standard, it will be indicated in the
annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality
standards - estimated error rate less than 1 per 10,000 bases.

FEATURES	Location/Qualifiers
source	1..98056
	/organism="Homo sapiens"
	/mol_type="genomic DNA"
	/db_xref="taxon:9606"
	/chromosome="X"
	/clone="GSHB-19024"
misc_feature	1..1599
	/notes="overlaps bases 151178..153177 of clone AC117405"
	/function="clone overlap"
repeat_region	40..82
	/rpt_family="L2"
repeat_region	378..507
	/rpt_family="L1MD3"
repeat_region	complement(518..570)
	/rpt_family="(GGA)n"
repeat_region	571..729
	/rpt_family="L1MC4"
repeat_region	730..987
	/rpt_family="MSTD"
repeat_region	complement(991..1400)
	/rpt_family="L2"
repeat_region	complement(1849..1779)
	/rpt_family="MIR"
repeat_region	complement(1783..2212)
	/rpt_family="LTR16A"
repeat_region	complement(3626..3656)
	/rpt_family="(GA)n"
repeat_region	complement(4713..5074)
	/rpt_family="THE1B"
repeat_region	5702..7108
	/rpt_family="L2"
repeat_region	complement(8670..8734)
	/rpt_family="MLTIG"
repeat_region	complement(9069..9144)
	/rpt_family="MLTIG"
repeat_region	complement(9254..9351)
	/rpt_family="MIR"
repeat_region	complement(9537..9688)
	/rpt_family="MER5A"
repeat_region	9797..9942
	/rpt_family="MIR"
repeat_region	complement(10403..10429)
	/rpt_family="AT-rich"
repeat_region	complement(10673..10705)
	/rpt_family="(CA)n"
repeat_region	complement(11460..11755)
	/rpt_family="AluSx"
repeat_region	1197..12125
	/rpt_family="MER5A"
repeat_region	12556..13643
	/rpt_family="L1MD2"
repeat_region	13618..13808
	/rpt_family="L1MD3"
repeat_region	complement(13997..14297)
	/rpt_family="MER4D"
repeat_region	14327..14625
	/rpt_family="AluSg"
repeat_region	complement(14648..14915)
	/rpt_family="MER4D"
repeat_region	complement(14915..15147)
	/rpt_family="MER4D"
repeat_region	15286..15404
	/rpt_family="FLAM_C"

JOURNAL

Submitted (14-FEB-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Apr 11, 2000 this sequence version replaced gi:6970579.
 All repeats were identified using RepeatMasker:
 Smith, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

COMMENT

----- Genome Center.
 Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR

Web site: <http://www-seq.wi.mit.edu>

Contact: sequence.submissions@genome.wi.mit.edu

----- Project Information

Center project name: L6731

Center clone name: 767 B.5

----- Summary Statistics

Sequencing vector: M13; M7815; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 185633 bases at least Q40

Consensus quality: 195115 bases at least Q30

Consensus quality: 201837 bases at least Q20

Insert size: 178000; agarose-fp

Quality coverage: 5.8 in Q20 bases; agarose-fp

Quality coverage: 4.9 in Q20 bases; sum-of-contigs

 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 31 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence.
 * as soon as it is available and the accession number will
 * be preserved.

```

1
* 1044: contig of 1043 bp in length
* 1044: gap of 100 bp
* 1144: contig of 1081 bp in length
* 2225: gap of 100 bp
* 2325: contig of 1021 bp in length
* 3345: gap of 100 bp
* 3445: contig of 1057 bp in length
* 4502: gap of 100 bp
* 4602: contig of 1154 bp in length
* 4903: gap of 100 bp
* 5757: gap of 100 bp
* 5857: contig of 1241 bp in length
* 7098: gap of 100 bp
* 7198: contig of 1039 bp in length
* 8237: gap of 100 bp
* 8337: contig of 1017 bp in length
* 9354: gap of 100 bp
* 9454: contig of 1252 bp in length
* 10706: gap of 100 bp
* 10806: contig of 1527 bp in length
* 12333: gap of 100 bp
* 12433: contig of 1445 bp in length
* 13878: gap of 100 bp
* 13978: contig of 1224 bp in length
* 15202: gap of 100 bp
* 15302: contig of 1667 bp in length
* 16968: gap of 100 bp
* 17069: contig of 1992 bp in length
* 19061: gap of 100 bp
* 19161: contig of 3479 bp in length
* 22639: gap of 100 bp
* 22640: contig of 3637 bp in length
* 26377: gap of 100 bp
* 26477: contig of 2799 bp in length
* 29375: gap of 100 bp
* 29376: contig of 3890 bp in length
* 33265: gap of 100 bp
* 33366: contig of 3647 bp in length
* 37013: gap of 100 bp
* 37113: contig of 2278 bp in length

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FEATURES
source

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* 39391: gap of 100 bp
* 39491: contig of 4200 bp in length
* 43690: gap of 100 bp
* 43790: contig of 3831 bp in length
* 47621: gap of 100 bp
* 47721: contig of 6131 bp in length
* 53852: gap of 100 bp
* 53952: contig of 9470 bp in length
* 63422: gap of 100 bp
* 63522: contig of 13839 bp in length
* 77361: gap of 100 bp
* 77461: contig of 15004 bp in length
* 92465: gap of 100 bp
* 92565: contig of 14386 bp in length
* 106951: gap of 100 bp
* 107051: contig of 22827 bp in length
* 129678: gap of 100 bp
* 129778: contig of 21649 bp in length
* 151427: gap of 100 bp
* 151527: contig of 30311 bp in length
* 181838: gap of 100 bp
* 181938: contig of 30182 bp in length
* 212120: contig of 30182 bp in length.

Location/Qualifiers
1. 212120
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="14"
/map="14"
/clone="RP11-767B5"
/clone_lib="RPC1-11 Human Male BAC"
1. 1043
/note="assembly_fragment"
1144. 2224
/note="assembly_fragment"
2325. 3345
/note="assembly_fragment"
3446. 4502
/note="assembly_fragment"
4603. 5756
/note="assembly_fragment"
5857. 7097
/note="assembly_fragment"
7198. 8236
/note="assembly_fragment"
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9454. 10705
/note="assembly_fragment"
10806. 12332
/note="assembly_fragment"
12433. 13877
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13978. 15201
/note="assembly_fragment"
15302. 16968
/note="assembly_fragment"
17069. 19060
/note="assembly_fragment"
19161. 22639
/note="assembly_fragment"
22740. 26376
/note="assembly_fragment"
26477. 29275
/note="assembly_fragment"
29376. 33265
/note="assembly_fragment"
33366. 37012
/note="assembly_fragment"
37113. 39390
/note="assembly_fragment
clone_end:SP6
vector_side:right"

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misc_feature

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Db 114062 GTTTTCAACTCTGATCCCGCATGGTTTTTATATATTCGTACACAAAGAGATTAG 114121
QY 612 ACAGTAAGAGTTTACAGAAAT-AAATCTATATTTTGTGAAGGTAGTGGTATTATAT 670
Db 114122 ACAGTAAGAGTTTACAGAAATAAATCTATATTTTGTGAAGGTAGTGGTACTAT 114181
QY 671 TGAGATTCAGTAGTTTCTAAGTCGTGTTATGTTTGTGTATACAAATGCGAGGTTTACACG 730
Db 114182 GTAGATTTTCAGTAGTTTCTAAGTCGTGTTATGTTTGTGTATACAAATGCGAGGTTTACACG 114241
QY 731 TCTATGCAATTTGACAAAAAGTTTAAAGAACTACATGTAATACTTGTAGTAAAT 790
Db 114242 TCTATGCAATTTGACAAAAAGTTTAAAGAAA--ACATGTAATACTTGTAGTAAAT 114298
QY 791 AACTTGCCATTTCTTATATGAAGCAGTATTTGGTGTGTTAAAAATTATAACAGTTAT 850
Db 114299 AACTTGCCATTTCTTATATGAAGCAGTATTTGGTGTGTTAAAAATTATAACAGTTAT 114358
QY 851 AAAGAAGATTTATAAAGGAA 871
Db 114359 AAAGAAGATTTATAAAGTAAACTAAA 114379

RESULT 5
AC108338 272578 bp DNA linear HTG 08-OCT-2002
LOCUS Rattus norvegicus clone CH230-114P1, *** SEQUENCING IN PROGRESS
DEFINITION ***; 5 unordered pieces.
ACCESSION AC108338
VERSION HTG108338.4 GI:23101239
KEYWORDS HTG; HTGS PHASE1; HTGS DRAFT; HTGS_ENRICHED.
SOURCE Rattus norvegicus (Norway rat)
ORGANISM Rattus norvegicus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 272578)
Muzny,D,Marie., Metzker,M.Lee., Abramzon,S., Adams,C., Alder,J.,
Allen,C., Allen,H., Alsbrooks,S., Amin,A., Anguiano,D.,
Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,
Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,F.,
Biswal,K., Blair,J., Blankenburg,K., Blyth,P., Brown,M.,
Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,
Cardenas,V., Carter,K., Cavazos,I., Ceasar,H., Center,A.,
Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Y., Chen,Z., Chu,J.,
Cleveland,C., Cockrell,R., Cox,C., Coyle,M., Cree,A., D'Souza,L.,
Davilla,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,
Delgado,O., Denson,S., Deramo,C., Ding,Y., Dinh,H., Divya,K.,
Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,
Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G.,
Fernandez,S., Finley,M., Flagg,N., Forbes,L., Foster,M., Foster,P.,
Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,
Gebregiorgis,E., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,W.,
Gunaratne,P., Haaland,W., Hamil,C., Hamilton,C., Hamilton,K.,
Harvey,Y., Hawlak,P., Hawes,A., Henderson,N., Hernandez,J.,
Hernandez,R., Hines,S., Hladun,S.L., Hodgson,A., Hogues,M.,
Hollins,B., Howells,S., Huiyik,S., Hume,J., Idlebird,D., Jackson,A.,
Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,
Karpach,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
Kowis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,
Lorensheua,L., Loulsegged,H., Lozado,R.J., Lu,X., Ma,J.,
Maheshwari,M., Maindarte,M., Mahmoud,M., Malloy,K., Mangum,A.,
Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E.,
Mawhiney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,
Milosavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,
Morgan,M., Morris,K., Morris,S., Munidasa,M., Murphy,S.,
Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S.,
Nwaokeme,O., Okwuonu,G., Olarnpunsagoon,A., Pal,S., Parks,K.,
Pasternak,S., Paul,H., Perez,A., Perez,L., Pfannkoch,C.,
Plopper,F., Poindexter,A., Popovic,D., Primus,E., Pu,L.-L.,
Puzo,M., Quiroz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R.,
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,

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Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J.,
Sanders,W., Savery,G., Scher,S., Scott,G., Shatsman,S., Shen,H.,
Shetty,J., Shvartbeyn,A., Sisson,I., Sitter,C.D., Sma's,D.,
Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Soza,J.,
Steinle,M., Strong,R., Sutton,A., Svatek,A., Taber,P., Taylor,C.,
Taylor,T., Thomas,N., Thomas,S., Tingle,A., Trejos,Z., Usmani,K.,
Valas,R., Vera,V., Villaseana,D., Waldron,L., Walker,B., Wang,J.,
Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,
Williams,G., Willson,R., Wlezyk,R., Woodson,H., Worley,K.,
Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,
Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von
Niederhausern,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs,R.A.
Direct Submission
Unpublished
2 (bases 1 to 272578)
Worley,K.C.
Direct Submission
Submitted (27-JAN-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 272578)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (08-OCT-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Sep 18, 2002 this sequence version replaced gi:21737649.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly ('contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GPMV
Center clone name: CH230-114P1
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 218776 bases at least Q40
Consensus quality: 22844 bases at least Q30
Consensus quality: 225163 bases at least Q20
Estimated insert size: 245370; sum-of-contigs estimation
Quality coverage: 4x in Q20 bases; sum-of-contigs estimation
* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 5 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 257855: contig of 257855 bp in length
* 257856: gap of unknown length
* 257956: contig of 3824 bp in length
* 261779: gap of unknown length
* 261879: gap of unknown length
* 261980: contig of 1235 bp in length
* 263114: gap of unknown length
* 263214: gap of unknown length
* 263215: contig of 2990 bp in length

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TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

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* 266205 266304: gap of unknown length
* 266305 272578: contig of 6274 bp in length.
FEATURES
    source
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            /mol_type="genomic DNA"
            /db_xref="taxon:10116"
            /clone="CH230-114P1"
        1..1828
            /notes="wgs contig"
        257956..259766
            /notes="wgs_contig"
    ORIGIN
        Query Match      43.3%; Score 431.4; DB 2; Length 272578;
        Best Local Similarity 80.5%; Pred. No. 9e-78;
        Mismatches 611; Conservative 0; Mismatches 121; Indels 27; Gaps 8;
    117 TGTAGAAATCTGTGCTTATCTATCGAATCTTTTGATATATATACATTGGGAA-CC 175
    Db 237426 TGTATAGAGCCTTATACCGCTCTATGAAGTCCCTTTGAGATGTTGTAGGAGGAATGC 237485
    QY 176 TGAATGTAGC---TTGACATTTTCCATGTAAACACAGTAGCCTGATCCACATTAAAC 232
    Db 237486 TGAATGCATCAATTACATTTTCCCATCAATGAGCCACGCTGATCCAGCATTAAGC 237545
    QY 233 TGTACTAACAACAACAGTGTATAGCTTCAATTAATGAAGCTTTGCTTTCTCTGGAAC 292
    Db 237546 TAATACCAAC-----TCCATGCAATGCTTCACTAAACAGGCTTTGCTGCTGTAGAAAT 237600
    QY 293 TGGTGAATAATCAAACTGTTGTGTACACCCCTGATGAGCTTCTGTTGCTCTTCCAC 352
    Db 237601 GGT---AAACTGGAGTGTGTTCTGATACCTTCAATGAGCTTCTGTTGCTTTTCC 237657
    QY 353 CAGAAATGGGGAATGATTTCCCAAATGGCAAGAAACAGAGTGATGCTATCTATCTGCAC 412
    Db 237658 ---AGAAATGAATGATTTCCCATTTGGCAACCA-----GGGCTACAATGCTGCAC 237706
    QY 413 CTTTGTGAAGTGTGCTTTCTTTCTTTGTTTCCAGGACACATGAGGAGTCTTT 472
    Db 237707 ACTTTGTAAGCTCTTTTCTTTTCTTTGTTT-CAGGACACAATGAGGAGGCTTT 237765
    QY 473 TCCACATGGCAGATGATTTGGGCAGAGCGATGAGTCCCTTAGTATCAGTCATGACAGATG 532
    Db 237766 TCCACATGGCAGATGATTTGGGCAGAGCGATGAGTCCCTTAGTTCAGTCATGATAGTG 237825
    QY 533 AAGAAGGAGCAGATAAATGTTTCAACTCTGATCCCGCATGTTTATATATATTC 592
    Db 237826 AAGAAGGAGCAGATAAATGTTTCAACTCTGATCCCTGACCCCGCATGTTTATATATATTC 237885
    QY 593 ATACAAAGAGAGATTAGACAGTAAAGATTACAGAAATAAATCTATATTTTGTGAA 652
    Db 237886 GTACACAAAGAGATTAGACAGTAAAGATTACAGAAATAAATCTATATTTTGTGAA 237945
    QY 653 GGGTAGTGGTATATACCTGATGATTTTCAAGTCTGATGTTTATGTTTGTAACT 712
    Db 237946 GGGTAGTGGTACTATACCTGATGATTTTCAAGTCTGATGTTTATGTTTGTAACT 238005
    QY 713 AATGGCAGGTTTACACGCTGATGCAATGCTACAAAAGATTATAGAAAACCTACATGA 772
    Db 238006 AATGGCAGGTTTACACGCTGATGCAATGCTACAAAAGATTATAGAAAACCTACATGA 238062
    QY 773 AAATCTGTAGTAAATAACTTGCATTTCTTTATATGAAACGCAATTTGGTGTGTTA 832
    Db 238063 AAATCTGTAGTAAATAACTTGCATTTCTTTATATGAAACGCAATTTGGTGTGTTA 238122
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LOCUS      AX817316
DEFINITION Sequence 64 from Patent WO02081517.
ACCESSION  AX817316
VERSION     AX817316.1 GI:39722703
KEYWORDS
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1
AUTHORS     Decristofaro,M.F., Padigaru,M., Miller,C., Tchernev,V., Zhong,H.,
            Zhong,M., Anderson,D., Ballinger,R., Gerlach,V., Spytek,K.A.,
            Rastelli,L., Kekuda,R., Guo,X., Zethusen,B., Andrew,D., Mezes,P.,
            Patturajan,M., Burgess,C.E., Eisen,A., Wolenc,A., Baumgartner,J.,
            Shimkets,R.A., Gusev,V., Vernet,C.A., Taupier,R.J., Pena,C.,
            Shenoy,S., Li,L., Casman,S., Bolgog,F., Fernandes,E., Smithson,G.,
            Malyankar,U., Tailon,B. and Liu,X.
TITLE       Novel polypeptides and nucleic acids encoded thereby
JOURNAL     Patent: WO 0208151-A 64 17-OCT-2002;
            Curagen Corporation (US)
FEATURES    Location/Qualifiers
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            41..44; Score 412.2; DB 6; Length 2463;
            Query Match      41.4%; Score 412.2; DB 6; Length 2463;
            Best Local Similarity 97.0%; Pred. No. 1.5e-73;
            Mismatches 420; Conservative 0; Mismatches 13; Indels 0; Gaps 0;
    QY 439 CTTTGTGTTTCCAGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGTTTGGGAGA 498
    Db 1998 CCTAGTTTCAAGAGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGTTTGGGAGA 2057
    QY 499 CGGATGGAGTCTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTAC 558
    Db 2058 CGGATGGAGTCTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTAC 2117
    QY 559 AACTCCTGATTCGCCGATGTTTATTAATATTCATACAAAGAGAGGATTAGACAGTAA 618
    Db 2118 AACTCCTGATTCGCCGATGTTTATTAATATTCATACAAAGAGAGGATTAGACAGTAA 2177
    QY 619 GAGTTTACAAGAAATAAATCTATATTTTGTGAAGGGTAGTGGTATTATCTAGTAGATT 678
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    QY 679 CAGTAGTTTCTAAGTCTGTTATTGTTTGTGTTTGAAGTGGAGGTTTACAGTCTATGCA 738
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    QY 739 ATTGTACAAAAAGTTATAAGAAAACACTACATGTAAAAATCTTGTAGCTAAATAACTTGCC 798
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799 ATTCTTTATATGGAACGATTTTGGTTGTTTAAAAATTTTAAACAGTTTATAAAGAAAG 858
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859 AATTATAAGGAA 871
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2418 ATTGTAACATAA 2430
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RESULT 7
BC028720 4658 bp mRNA linear PRI 25-AUG-2003
LOCUS Homo sapiens dystrophin (muscular dystrophy, Duchenne and Becker
DEFINITION types), transcript variant Dp71b, mRNA (CDNA clone IMAGE:482807),
complete cds.
ACCESSION BC028720
VERSION BC028720
KEYWORDS BC028720.1 GI:20379675
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
MAMMALIA; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 4658)
AUTHORS Strausberg,R.L., Feingold,E.A., Grouse,L.H., Derge,J.G.,
Klausner,R.D., Collins,F.S., Wagner,L., Shenmen,C.M., Schuler,G.D.,
Altschul,S.F., Zeeberg,B., Buetow,K.H., Schaefer,C.F., Bhat,N.K.,
Hopkins,R.F., Jordan,H., Moore,T., Max,S.I., Wang,J., Hsieh,P.,
Diatchenko,K., Marusina,K., Farmer,A.A., Rubin,G.M., Hong,L.,
Stapleton,T.E., Soares,M.B., Bonaldo,M.F., Casavant,T.L.,
Schetz,T.M., Brownstein,M.J., Udwin,T.B., Toshiyuki,S.,
Carninci,P., Prange,C., Raha,S.S., Loguillano,N.A., Peters,G.J.,
Abramson,R.D., Mullahy,S.J., Bosak,S.A., McEwan,P.J.,
McKernan,K.J., Malek,J.A., Gunaratne,P.H., Richards,S.,
Worley,K.C., Hale,S., Garcia,A.M., Gay,L.J., Hulyk,S.W.,
Vallalao,D.K., Muzny,D.M., Sodergren,E.J., Lu,X., Gibbs,R.A.,
Fahey,J., Heiton,E., Kettelman,M., Madan,A., Rodrigues,S.,
Sanchez,A., Whiting,M., Madan,A., Young,A.C., Shevchenko,Y.,
Bouffard,G.G., Blakesley,R.W., Touchman,J.W., Green,E.D.,
Dickson,M.C., Rodriguez,A.C., Grimwood,J., Schmutz,J., Myers,R.M.,
Butterfield,Y.S., Krzywinski,M.I., Skalska,U., Smalish,D.E.,
Schnerch,A., Schein,J.E., Jones,S.J. and Marra,M.A.
Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences
Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
22388257
12477932
2 (bases 1 to 4658)
Strausberg,R.
Direct Submission
Submitted (29-APR-2002) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA
REMARK NTH-MGC Project URL: http://mgc.nci.nih.gov
COMMENT Contact: MGC help desk
Email: cgaps@mail.nih.gov
Tissue Procurement: Miklos Falkovits, M.D., Ph.D.
cDNA Library Preparation: Michael J. Brownstein (NHGRI) & Shiraki
Toshiyuki and Piero Carninci (RIKEN)
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Institute for Systems Biology
http://www.systemsbio.org
contact: amadan@systemsbio.org
Anup Madan, Jessica Fahey, Erin Helton, Mark Kettelman, Anuradha
Madan, Stephanie Rodrigues, Amy Sanchez and Michelle Whiting
Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
Series: IRAP Plate: 46 Row: 1 Column: 3
This clone was selected for full length sequencing because it
passed the following selection criteria: matched mRNA gi: 503296
This clone has the following problem: The cds is short compared to
the longest cds in the locus.

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FEATURES

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Location/Qualifiers

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 RVAKGHQHYPMVEYCTPTSGEDVRDFAKVLNKKERTKRYFAKHPRMGYLPQVLE
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misc_feature

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 calmodulin Putative zinc finger"
 /db_xref="CDD:pfam00569"

ORIGIN

Query Match 41.4%; Score 412.2; DB 9; Length 4658;
 Best Local Similarity 97.0%; Pred. No. 1.3e-73;
 Matches 420; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

QY 439 CTTTGTTCAGGACACAAATGTAGGAAGTCTTTTCACATGGCAGATGTTGGGCGA 498
 DB 1927 CCTAGTTCAAGAGGACAAATGTAGGAAGTCTTTTCACATGGCAGATGTTGGGCGA 1986

QY 499 CGATGGAGTCTTGTAGTATCAGTCATGACAGATGACAGAGGAGAGATAAATGTTTAC 558
 DB 1987 CGATGGAGTCTTGTAGTATCAGTCATGACAGATGACAGAGGAGAGATAAATGTTTAC 2046

QY 559 AACTCCTGATCCCGCATGTTTATAATATTCATACAAAGAGAGGATTAGACAGTAA 618
 DB 2047 AACTCCTGATCCCGCATGTTTATAATATTCATACAAAGAGGATTAGACAGTAA 2106

QY 619 GAGTTTACAGAAATAATCTATATTTTGTGAAGGAGTGGTATTATCTAGTAGATT 678
 DB 2107 GAGTTTACAGAAATAATCTATATTTTGTGAAGGAGTGGTATTATCTAGTAGATT 2166

QY 679 CAGTAGTTTCTAAGTCTGTTATGTTTGTAACTGGCAGGTTTACACGCTCTATGCA 738
 DB 2167 CAGTAGTTTCTAAGTCTGTTATGTTTGTAACTGGCAGGTTTACACGCTCTATGCA 2226

QY 739 ATTGTACAAAAGATTAAAGAAACATCATGTAAAACTCTTGTAGTAGCTAAATACTTGC 798
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QY 799 ATTCTTTATATGGAACGATTTTGGTTGTTTAAAAATTTTAAACAGTTTATAAAGAAAG 858
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Db      2347 ATTGTAACATAA 2359

RESULT 8
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DEFINITION Sequence 40 from Patent WO0229056.
ACCESSION AX538620
VERSION    AX538620.1 GI:252711166
KEYWORDS   synthetic construct
SOURCE     synthetic construct
ORGANISM   artificial sequences.

REFERENCE 1
AUTHORS    Chamberlain,J.S. and Harper,S.Q.
TITLE      Mini-dystrophin nucleic acid and peptide sequences
JOURNAL    Patent: WO 0229056-A 40 11-APR-2002;
           THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)
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Best Local Similarity 98.3%; Pred.No. 2.5e-73;
Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

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QY      570 CCGCATGGTTTATATAATTCATACAAAGAGGATTAGACAGTAAAGTTTACAAG 629
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QY      870 AA 871
Db      4911 AA 4912

RESULT 9
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LOCUS      AX538619
DEFINITION Sequence 39 from Patent WO0229056.
ACCESSION AX538619
VERSION    AX538619.1 GI:252711163
KEYWORDS   synthetic construct
SOURCE     synthetic construct

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ORGANISM   synthetic construct
REFERENCE 1
AUTHORS    Chamberlain,J.S. and Harper,S.Q.
TITLE      Mini-dystrophin nucleic acid and peptide sequences
JOURNAL    Patent: WO 0229056-A 39 11-APR-2002;
           THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)
FEATURES   Location/Qualifiers
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Best Local Similarity 98.3%; Pred.No. 2.5e-73;
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Db      4989 AA 4990

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LOCUS      AX538621
DEFINITION Sequence 41 from Patent WO0229056.
ACCESSION AX538621
VERSION    AX538621.1 GI:252711168
KEYWORDS   synthetic construct
SOURCE     synthetic construct
ORGANISM   artificial sequences.

REFERENCE 1
AUTHORS    Chamberlain,J.S. and Harper,S.Q.
TITLE      Mini-dystrophin nucleic acid and peptide sequences
JOURNAL    Patent: WO 0229056-A 41 11-APR-2002;
           THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)
FEATURES   Location/Qualifiers
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QY 870 AA 871
Db 9159 AA 9160

RESULT 13
LOCUS AX538627 12057 bp DNA linear PAT 23-NOV-2002
DEFINITION Sequence 47 from Patent WO0229056.
ACCESSION AX538627
VERSION AX538627.1 GI:25271181
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
          artificial sequences.
REFERENCE
  1. Chamberlain, J.S. and Harper, S.Q.
  Mini-dystrophin nucleic acid and peptide sequences
  Patent: WO 0229056-A 47 11-APR-2002;
  THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)
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ORIGIN
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Best Local Similarity 98.3%; Pred. No. 2.2e-73;
Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
QY 450 AGGACACAATGTAGGAAGTCTTTTCCACATGCGCAGATGATTTGGCAGAGCGATGAGTC 509
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Db 11634 AA 11635
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LOCUS HSDMDR 12446 bp mRNA linear PRI 12-SEP-1993
DEFINITION Human mRNA for dystrophin.
ACCESSION X14298
VERSION X14298.1 GI:30845
KEYWORDS Dmd gene; Duchenne muscular dystrophy; dystrophin.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
          1 (bases 1 to 12446)
          Rosenthal, A., Speer, A., Billwitz, H., Cross, G.S., Forrest, S.M. and
          Davies, K.E.
          Two human cDNA molecules coding for the Duchenne muscular dystrophy
          (DMD) locus are highly homologous
          Nucleic Acids Res. 17 (13), 5391 (1989)
          89345106
          266885
          2 (bases 1 to 12446)
          Rosenthal, A.
          Direct Submission
          Submitted (09-FEB-1989) Rosenthal A., Akademie der Wissenschaften
          der DDR, Zentralinstitut fuer Molekularbiologie, Robert-Roesle
          Str.10, 1115 Berlin Buch, DDR
          See also M18533 and M20250 for Dmd segs.; discrepancies compared to
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            ETVTITTRQILVKHAQEBELPPPKPKKQITVDSEIRKLDVDITELHSLTRSEA
            VLOSPEFAI PRKGNFSDLEKVNALEREKAFKRLQDASRSGQALVEGMVNEGVA
            DSIQASEQLNSRWIEFCQLLSERLWLEYQNNIIAPYNQLQLEQMTTAEWNLKIQ
            PTPSEPTATKSQLKICKEVNRLSLQPOIERLKIOSIALKEKGQGMFLDADFAF
            TNHFQVSDVQAREKELQITFDLPMPYQETMSAIRTWQOSETKLSIPQSLVDY
            EIMORLGEIQALQSLQEQSGLYLSTTVKEMSKAPSEISRKYSQSEFEIEGRWK
            KLSQIVHEQKLEBEOMNKLQNLQTLKWMAEVDVFLKEWPAKGSEILKKOL
            KQCLLVSDITQTPSLNSVNEGQKIKNEAEPEFASRLTELKELNTDQHMCOQVY
            ARKALGGKLEKTVSLQDUSEHEMWTQAESEYLELDEFEYKTFDELQAKFEEMKRAK
            EEAQKRAKVLKTESVNSVIAQPPVAQEAALKKELETTITNTQMLCTRLNGKKTLE
            EVWACHMELLSYLEKANKNLNEVEFLKTTENIPGGABEISEVLDLSLENLRHSEDNF
            NQIRIACITLTDGWMDELNEELTFNSRWELHEEAVRQKLEQSIQSAQETNS
            LHLQESLFTDKOLAAVIAQVDAQPOAKIQSDLSHETSLSEMKKHNGKGA
            AQVLSQIDVACKQLQDVSMKFLFOKPAFNEQFLOESKMLDVKHMLPALETKSYE
            QEVQSLNHCNLYKSLSEVSEVMVITGQIVQKQTEPNKELDERVTKLHY
            NELGAKVTERKQKLEKLUKSRKEMNVITELAAATDMELTFKSAVEGMPNLDSE
            VAWGATQRETEKQKHLKSI TEVGEALKTVLGKKEITVLELAATDMELTFKSAVEGMPNLDSE
            EWLNLLEYQKHEMTFDQNVDHITKWIQADTLLDSESEKKKPKQKEDVLRKLKALND
```

[illegible]

GB2368064-A.
24-APR-2002.
16-JAN-2001; 2001GB-00001124.
30-SEP-2000; 2000US-0237079P.
(IMCR) IMPERIAL CANCER RES TECHNOLOGY LTD.
(BARB/) BARBER E.
Barber E;
WPI; 2002-429042/46.
P-PSDB; AAU98739.
New human regulatory polynucleotide, useful for treating disorders associated with protein truncation, particularly muscular dystrophy, and related peptides and antibodies.
Claim 5; Page 170-172; 222pp; English.
The invention relates to a polynucleotide (I) comprising, or consisting of, apo-dystrophin-4 inversion sequence appearing as ASK86496, or its functional equivalents (e.g. the apo-dystrophin-4 cDNA sequence appearing as ASK86497). Also included are polynucleotides that hybridise to either strand of (I), a vector containing (I), a cell containing (I) or the vector, proteins and peptides encoded by (I), a protein homologous with human dystrophin that is expressed on cell surfaces in vivo antibodies (Ab) specific for the protein and method of screening for leukemia cells by analysing DNA for the presence of (I) or by detecting presence of (II). The apo-dystrophin-4 inversion sequence is a regulatory element that controls expression (transcription and translation) of associated DNA, and may allow read-through of stop codons. The apo-dystrophin-4 inversion sequence is used in gene therapy of diseases associated with truncation of proteins, particularly muscular dystrophy and also leukaemia, but more generally (i) is a regulatory sequence used to control expression of any attached gene. Analysis of DNA for (I), or detection of proteins (II) encoded by (I), can be used to screen for leukaemic cells and related diseases. Antibodies raised against (II) can be used therapeutically, to inhibit (II) activity, also to detect (II) in screening assays. The present sequence is the cDNA sequence for human apo-dystrophin-4 containing a plurality of stop codons some of which may be read through due to the presence of (I) in the apo-dystrophin-4 gene
Sequence 996 BP; 334 A; 152 C; 195 G; 315 T; 0 U; 0 Other;
Query Match 100.0%; Score 996; DB 5; Length 996;
Best Local Similarity 100.0%; Pred. No. 2.4e-211;
Matches 996; Conservative 0; Mismatches 0; Indels 0; Gaps 0
QY 1 GTGGTTTGATTGATAGTAAAAAATGTTTCGTTAATCAAGTAGAGAGTAAGTAATCAAT 60
Db 1 GTGGTTTGATTGATAGTAAAAAATGTTTCGTTAATCAAGTAGAGAGTAAGTAATCAAT 60
QY 61 CAATCACTCATGACCAAGTGGAAGAAGATGTATCCCATCATGGAATATTCCTGTTCTGAT 120
Db 61 CAATCACTCATGACCAAGTGGAAGAAGATGTATCCCATCATGGAATATTCCTGTTCTGAT 120
QY 121 AGAATACTTGTCCTTATCTATGGAATCTTTTGATATATATTACATTTGGGAACCTGAAT 180
Db 121 AGAATACTTGTCCTTATCTATGGAATCTTTTGATATATATTACATTTGGGAACCTGAAT 180
QY 181 GTAGCTTGACATTTTCCATGTAACACACCATGACGCTGATCCCAACATTAAGCTGATACATA 240
Db 181 GTAGCTTGACATTTTCCATGTAACACACCATGACGCTGATCCCAACATTAAGCTGATACATA 240
QY 241 ACAACAACAGCTGTAATGGCTTCATTAAAGAGCTTTTCCTCTCTCGGAACCTGGTGA 300
Db 241 ACAACAACAGCTGTAATGGCTTCATTAAAGAGCTTTTCCTCTCTCGGAACCTGGTGA 300
QY 301 AATCAAAACCTTGTTGTGACACCCCTCGATGCGAGCTCTGTGTGTGTCTTCCACCAGAAATG 360

FT FT /transl_except= .453,aa:Xaa) .453,aa:Xaa)
FT FT /transl_except= .489,aa:Xaa) .489,aa:Xaa)
FT FT /transl_except= .504,aa:Xaa-Xaa) .504,aa:Xaa-Xaa)
FT FT /transl_except= .630,aa:Xaa) .630,aa:Xaa)
FT FT /transl_except= .781,aa:Xaa) .781,aa:Xaa)
FT FT /transl_except= .801,aa:Xaa) .801,aa:Xaa)
FT FT /transl_except= .852,aa:Xaa) .852,aa:Xaa)
FT FT /transl_except= .885,aa:Xaa) .885,aa:Xaa)
FT FT /transl_except= .920,aa:Xaa) .920,aa:Xaa)
FT FT /transl_except= .947,aa:Xaa) .947,aa:Xaa)
FT FT /transl_except= .1026,aa:Xaa) .1026,aa:Xaa)
FT FT /transl_except= .1075,aa:Xaa) .1075,aa:Xaa)
FT FT /transl_except= .1086,aa:Xaa) .1086,aa:Xaa)
FT FT /transl_except= .1114,aa:Xaa) .1114,aa:Xaa)
FT FT /transl_except= .1143,aa:Xaa) .1143,aa:Xaa)
FT FT /transl_except= .1188,aa:Xaa) .1188,aa:Xaa)
FT FT /transl_except= .1216,aa:Xaa) .1216,aa:Xaa)
FT FT /transl_except= .1224,aa:Xaa) .1224,aa:Xaa)
FT FT /note= "Xaa= unknown, encoded by in frame stop codon"
PN GB2368064-A.
XX
XX
XX PD 24-APR-2002.
XX
XX 16-JAN-2001; 2001GB-00001124.
XX
XX 30-SEP-2000; 2000US-0237079P.
PR
PR (IMCR) IMPERIAL CANCER RES TECHNOLOGY LTD.
PA (BARB/) BARBER E.
PA
XX
XX Barber E;
PI
PI WPI: 2002-429042/46.
DR P-PSDB; AAU98729.
DR
XX
XX New human regulatory polynucleotide, useful for treating disorders
PT associated with protein truncation, particularly muscular dystrophy, and
PT related peptides and antibodies.
XX
XX Disclosure; Fig 4; 222pp; English.
PS
XX
XX The invention relates to a polynucleotide (I) comprising, or consisting
CC of, apo-dystrophin-4 inversion sequence appearing as ABK86496, or its
CC functional equivalents (e.g. the apo-dystrophin-4 cDNA sequence appearing
CC as ABK86497). Also included are polynucleotides that hybridize to either
CC strand of (I), a vector containing (I), a cell containing (I) or the
CC vector, proteins and peptides encoded by (I), a protein homologous with
CC human dystrophin that is expressed on cell surfaces in vivo antibodies
CC (Ab) specific for the protein and method of screening for leukemia cells
CC by analysing DNA for presence of (I) or by detecting presence of (I).
CC The apo-dystrophin-4 inversion sequence is a regulatory element of (I).
CC controls expression (transcription and translation) of associated DNA,
CC and may allow read-through of stop codons. The apo-dystrophin-4 inversion
CC sequence is used in gene therapy of diseases associated with truncation
CC of proteins, particularly muscular dystrophy and also leukaemia, but more
CC generally (I) is a regulatory sequence used to control expression of any
CC attached gene. Analysis of DNA for (I), or detection of proteins (II)
CC encoded by (I), can be used to screen for leukemic cells and related
CC diseases. Antibodies raised against (II) can be used therapeutically, to
CC inhibit (II) activity, also to detect (II) in screening assays. The
CC present sequence is the full length cDNA sequence for human apo-
CC dystrophin-4 containing a plurality of stop codons some of which may be
CC read through due to the presence of (I) in the apo-dystrophin-4 gene
XX
XX Sequence 1230 BP; 404 A; 189 C; 259 G; 378 T; 0 U; 0 Other:

```
Query Match      100.0%; Score 996; DB 6; Length 1230;
Best Local Similarity 100.0%; Pred. No. 2.5e-211;
Matches 996; Conservative 0; Mismatches 0; Indels 0; Gaps 0
```

QY 1 GTGGTTTGATTGATAGTAAAAAATGTTCTTAATACAAGTAGAGACTAAGTAATCAAT 60

Db	235	GTGGTTTGATGTGATGATAAAAAAATGTTCCGTTAATAACAAGTAGAGAGTAAGTAATCAAT	294
Qy	61	CAATCACTCATAGCCAAAGTGGAAGAAGATGTATCCCATCATGGAATATTCCTGTTCTGAT	120
Db	295	CAATCACTCATAGCCAAAGTGGAAPAGATGTATCCCATCATGGAAATATTCCTGTTCTGAT	354
Qy	121	AGAAATCTTGTCCTTATCTATGGAATCTTTTGATATATATTTACATTTGGAAACCTGAAAT	180
Db	355	AGAAATCTTGTCCTTATCTATGGAATCTTTTGATATATATTTACATTTGGAAACCTGAAAT	414
Qy	181	GTAGCTTGACATTTTTCCTATGTAAACACCAAGTAGCCTGATCCAAACATTAAGCTGTACTTA	240
Db	415	GTAGCTTGACATTTTTCCTATGTAAACACCAAGTAGCCTGATCCAAACATTAAGCTGTACTTA	474
Qy	241	ACAAAACAAAGTGTAATGGCTTCATTAATAAGGCTTTGCTTTCTTCCTGGAAAACGTGGTGA	300
Db	475	ACAAAACAAAGTGTAATGGCTTCATTAATAAGGCTTTGCTTTCTTCCTGGAAAACGTGGTGA	534
Qy	301	AATCAAAACCTTGTTGTGTACACCCTCGATGACGCTTCTGTTGTGCTCTCACCCAGAAAATG	360
Db	535	AATCAAAACCTTGTTGTGTACACCCTCGATGACGCTTCTGTTGTGCTCTCACCCAGAAAATG	594
Qy	361	GGGAATGATTTCCAAATGGCAAGAAACAGAGTGATCTATCTATCTGCACCTTTTGTA	420
Db	595	GGGAATGATTTCCAAATGGCAAGAAACAGAGTGATCTATCTATCTGCACCTTTTGTA	654
Qy	421	AAGCTGTCTTCTTCTCTTCTTTTCCAGNACAATGTAGGAAGTCTTTTCCACATG	480
Db	655	AAGCTGTCTTCTTCTCTTCTTTTCCAGNACAATGTAGGAAGTCTTTTCCACATG	714
Qy	481	GCAGATGATTTGGGCAGAGCGATGGAGTCTCTTAGTATCAGTCAATGACAGATGAAGAAGGA	540
Db	715	GCAGATGATTTGGGCAGAGCGATGGAGTCTCTTAGTATCAGTCAATGACAGATGAAGAAGGA	774
Qy	541	GCAGATTAATGTTTACAACTCCTGATTCGCGCATGGTTTTTAATATTCATACAACA	600
Db	775	GCAGATTAATGTTTACAACTCCTGATTCGCGCATGGTTTTTAATATTCATACAACA	834
Qy	601	AAGAGGATTAGACAGTAAGAGTTTACAAGAAATAAATCTATATTTTTGTGAAGGGTAGTG	660
Db	835	AAGAGGATTAGACAGTAAGAGTTTACAAGAAATAAATCTATATTTTTGTGAAGGGTAGTG	894
Qy	661	GTATTATACGTAGATTTCAAGTAGTTTCTAAGTCTCTTATGTTTGTTAACAATGGCAG	720
Db	895	GTATTATACGTAGATTTCAAGTAGTTTCTAAGTCTCTTATGTTTGTTAACAATGGCAG	954
Qy	721	GTTTTACAGTCTATGCAATGTPACAAAAGTTATAAGAAACACATCATGTAATCTCTG	780
Db	955	GTTTTACAGTCTATGCAATGTPACAAAAGTTATAAGAAACACATCATGTAATCTCTG	1014
Qy	781	ATAGCTAAATAACTTGCCATTTCTTATATGGAACGCATTTTGGGTGTTTAAAAATTTA	840
Db	1015	ATAGCTAAATAACTTGCCATTTCTTATATGGAACGCATTTTGGGTGTTTAAAAATTTA	1074
Qy	841	TACAGTTATAAGAAAGAAATATAAGGAAGAAAGAAATACGCAATGGACAAGTGGTG	900
Db	1075	TACAGTTATAAGAAAGAAATATAAGGAAGAAAGAAATACGCAATGGACAAGTGGTG	1134
Qy	901	AAGCTGTGAACCTCAGGTGTGCACAAATATCAGGAACACCCCAAAACCAAAAGTCAGGTAGA	960
Db	1135	AAGCTGTGAACCTCAGGTGTGCACAAATATCAGGAACACCCCAAAACCAAAAGTCAGGTAGA	1194
Qy	961	AATAGCATGAGAAGCCGTGTTGATGTTAAATTT	996
Db	1195	AATAGCATGAGAAGCCGTGTTGATGTTAAATTT	1230

RESULT 3

ABK86463

ID ABK86463 standard; cDNA; 1234 BP.

XX AC

XX ABK86463;

XX

PF 16-JAN-2001; 2001GB-00001124.
 PR 30-SEP-2000; 2000US-0237079P.
 XX (IMCR) IMPERIAL CANCER RES TECHNOLOGY LTD.
 PA (BARB/) BARBER E.
 XX Barber E;
 FI
 XX WP1; 2002-429042/46.
 DR
 XX
 PT New human regulatory polynucleotide, useful for treating disorders
 PT associated with protein truncation, particularly muscular dystrophy, and
 PT related peptides and antibodies.
 XX
 PS Disclosure; Fig 9; 222pp; English.
 XX
 CC The invention relates to a polynucleotide (I) comprising, or consisting
 CC of, apo-dystrophin-4 inversion sequence appearing as ABK86496, or its
 CC functional equivalents (e.g. the apo-dystrophin-4 cDNA sequence appearing
 CC as ABK86497). Also included are polynucleotides that hybridise to either
 CC strand of (I), a vector containing (I), a cell containing (I) or the
 CC vector, proteins and peptides encoded by (I), a protein homologous with
 CC human dystrophin that is expressed on cell surfaces in vivo antibodies
 CC (Ab) specific for the protein and method of screening for leukemia cells
 CC by analysing DNA for presence of (I) or by detecting presence of (II).
 CC The apo-dystrophin-4 inversion sequence is a regulatory element that
 CC controls expression (transcription and translation) of associated DNA,
 CC and may allow read-through of stop codons. The apo-dystrophin-4 inversion
 CC sequence is used in gene therapy of diseases associated with truncation
 CC of proteins, particularly muscular dystrophy and also leukaemia, but more
 CC generally (II) is a regulatory sequence used to control expression of any
 CC attached gene. Analysis of DNA for (I), or detection of proteins (II)
 CC encoded by (I), can be used to screen for leukaemic cells and related
 CC diseases. Antibodies raised against (II) can be used therapeutically, to
 CC inhibit (II) activity, also to detect (II) in screening assays. The
 CC present sequence is a region of mouse genomic DNA in the Apo-dystrophin-4
 CC gene region
 XX
 SQ Sequence 1044 BP; 347 A; 177 C; 200 G; 320 T; 0 U; 0 Other;

Query Match 43.7%; Score 435; DB 6; Length 1044;
 Best Local Similarity 84.7%; Fred. No. 7.1e-87;
 Matches 577; Conservative 0; Mismatches 80; Indels 24; Gaps 7;

QY 192 TTTTCCATGTAACACACAGTAGCTGATCCAACTTAAGCTGATCTAACAACAACGT 251
 DB 257 TTTCTCCCATCAATGACACCATGCTGATCCAGTATTAAAGCTAATACTAAC- 311
 QY 252 GTAATGGCTTCATTAAGGCTTGTCTTCTCTCGAACTGCTGAAATCAAACTT 311
 DB 312 TGCAATGCTTCATTAACAAGGATTGCTTCTTGTCTAGAAATGGGTAAAAA-CGGACTGT 369
 QY 312 GTTGTGTACACCTCGATGCGACTTCTGTGTGTCTTCCACCAAGAAATGGGAATGATTT 371
 DB 370 GGTCTGTATACCTTCAATGCGACTTATGTGTGTCTTTTCC--TGAAATGGTATGACTC 427
 QY 372 CCCAAATGGCAAGAAACAGAGTGTATGTATCTGCACTTTTGAAGTCTGTCTT 431
 DB 428 CCAATAGTGGCAACACAGGGGTACAACT- 476
 QY 432 TCTTTCTCTTTGTTTCCAGACACAATGAGGAGTCTTTCCACATGGCAGATGATTT 491
 DB 477 TCTTTCTCTTTGTTTCCAGACACAATGAGGAGTCTTTCCACATGGCAGATGATTT 536
 QY 492 GGGCAGAGCGATGGAGTCTTGTATGATGATGATGATGATGATGATGATGATGAT 551
 DB 537 GGGCAGAGCGATGGAGTCTTGTATGATGATGATGATGATGATGATGATGATGAT 596
 QY 552 GTTTTACAACTCTGATCCCGCATGGTCTTATATATTCATATTCATATTCATATTCAT 611
 DB 597 GTTTTACAACTCTGATCCCGCATGGTCTTATATATTCATATTCATATTCATATTCAT 656

QY 612 ACAGTAGAGTTTACAGAAAT-AAATCTATATTTTGTGAGGGTAGTGGTATATATACT 670
 DB 657 ACAGTAGAGTTTACAGAAATATAATTTTGTGAGGGTAGTGGTACTATATACT 716
 QY 671 GTAGATTTTCAAGTAGTTTCTAAGTCTGTATTGTTTGTAACTGGCAGGTTTACACG 730
 DB 717 GTAGATTTTCAAGTAGTTTCTAAGTCTGTATTGTTTGTAACTGGCAGGTTTACACG 776
 QY 731 TCTATGCAATTTGTACAAAAAGTTTATAGAAAACATCATATTAATCTTGATAGCTAAT 790
 DB 777 TCTATGCAATTTGTACAAAAAGTTTATAGAAAACATCATATTAATCTTGATAGCTAAT 833
 QY 791 AACTTGGCAATTTCTTATATGGAACGCAATTTTGGGTGTTTAAAAAATTTATAACAGTTAT 850
 DB 834 AACTTGGCAATTTCTTATATGGAACGCAATTTTGGGTGTTTAAAAAATTTATAACAGTTAT 893
 QY 851 AAAGAAAGAAATTTATAAGGAA 871
 DB 894 AAAGAAAGAAATTTATAAGGAA 914

RESULT 5
 ABT33375
 ID ABT33375 standard; DNA; 2463 BP.
 XX
 AC ABT33375;
 XX
 DT 22-MAY-2003 (first entry)
 XX
 DE NOVX DNA sequence SEQ ID No 64.
 XX
 KW Hepatotrophic; immunosuppressive; cardiant; hypertensive; tranquilizer;
 KW vulnaray; virucide; antibacterial; protozoacide; fungicide; nootropic;
 KW antiparasitic; neuroprotective; cerebroprotective; antiparkinsonian;
 KW anticonvulsant; antidiabetic; analgesic; dermatological; keratolytic;
 KW antiseborrheic; antihematuric; antiarthritic; antiinflammatory; anti-HIV;
 KW cytostatic; antischismatic; antipsoriatic; hypotensive; osteopathic;
 KW antitumor; anorectic; antidiabetic; antilept; antiallergic; naemostatic;
 KW neuroleptic; antidepressant; antiinfertility; NOVX; human disease;
 KW NOVX-associated disorder; trauma; viral; bacterial; fungal; protozoal;
 KW parasitic infection; Alzheimer's disease; stroke; forensic biology;
 KW immunogen; non-human transgenic animal; gene therapy; gene; ds.
 XX Unidentified.
 XX
 PN WO200281517-A2.
 XX
 PD 17-OCT-2002.
 XX
 XX 22-JAN-2002; 2002NO-US002064.
 PF
 XX 19-JAN-2001; 2001US-0262892P.
 PR 23-JAN-2001; 2001US-0263598P.
 PR 24-JAN-2001; 2001US-0263799P.
 PR 25-JAN-2001; 2001US-026417P.
 PR 26-JAN-2001; 2001US-0264139P.
 PR 26-JAN-2001; 2001US-0264478P.
 PR 30-JAN-2001; 2001US-0263351P.
 PR 02-MAR-2001; 2001US-0272870P.
 PR 14-MAR-2001; 2001US-0275927P.
 PR 15-MAR-2001; 2001US-0275990P.
 PR 20-MAR-2001; 2001US-0276449P.
 PR 20-MAR-2001; 2001US-0277358P.
 PR 23-MAR-2001; 2001US-0278151P.
 PR 23-MAR-2001; 2001US-0279857P.
 PR 20-APR-2001; 2001US-0285140P.
 PR 20-APR-2001; 2001US-0285141P.
 PR 30-APR-2001; 2001US-0287484P.
 PR 17-MAY-2001; 2001US-0291701P.
 PR 08-JUN-2001; 2001US-0296960P.
 PR 10-JUL-2001; 2001US-0304353P.
 PR 10-JUL-2001; 2001US-0304353P.
 PR 12-JUL-2001; 2001US-0304886P.

09-AUG-2001; 2001US-0311289P.
13-AUG-2001; 2001US-0311975P.
16-AUG-2001; 2001US-0312937P.
18-OCT-2001; 2001US-0330227P.
29-NOV-2001; 2001US-0334198P.
(CURA-) CURAGEN CORP.
Decristofaro MF, Padigaru M, Miller C, Tchernev V, Zhong H;
Zhong M, Anderson D, Ballinger R, Gerlach V, Sytek KA, Rastelli L;
Kekuda R, Guo X, Zerhusen B, Andrew D, Mezes P, Patturajan M;
Burgess CE, Eissen A, Wolenc A, Baumgartner J, Shimkets RA, Gusev V;
Vernet CAM, Taupier RJ, Pena C, Shenoy S, Li L, Casman S, Boldog F;
Fernandes E, Smithson G, Malyankar U, Taillon B, Liu X;
WPI: 2003-058504/05.
P-PSDB; AB373939.
New polypeptides, designated as NOVX, useful for diagnosing and treating
infections, neurological diseases, cancer, allergy, and bone,
immunological, skin, renal, brain, muscle and autoimmune disorders.
Claim 9; Page 171; 672pp; English.
The invention relates to a novel isolated polypeptide, designated NOVX
(NOVI - 33), consisting of a mature form of one of si sequences, given in
the specification, or its variant, where amino acid residue(s) in the
variant differ from the mature form, provided that the variant differs in
not more than 15 % of the amino acids from the sequence of the mature
form. The NOVX polypeptides, nucleic acids encoding the polypeptides, and
an antibody to the polypeptides, are useful for treating or preventing a
NOVX-associated disorder in humans and for treating a syndrome associated
with a human disease (NOVX-associated disorder). NOVX polypeptides and
the encoding nucleic acids, are useful for determining the presence of or
predisposition to a disease associated with altered levels of NOVX
polypeptide and polynucleotide, by measuring the level of polypeptide,
expression or the amount of nucleic acid from a mammal and comparing it
with another mammal not having or not predisposed to the disease. NOVX
polypeptide is also useful for identifying an agent that binds to NOVX
and a cell expressing NOVX is useful for identifying an agent that
modulates the expression or activity of NOVX. The antibodies and a
polypeptide having 95 % sequence identity to NOVX polypeptide are useful
for treating a pathological state in a mammal. The antibodies are also
useful for determining the presence or amount of NOVX in a sample. NOVX
polypeptides, polynucleotides and antibodies specific for the
polypeptides are useful for treating or preventing disorders or syndromes
including trauma, viral, bacterial, fungal, protozoal, and parasitic
infections. They can also treat disorders such as e.g., Alzheimer's
disease or a stroke. The NOVX encoding nucleic acids are useful for
expressing the NOVX proteins, to detect NOVX mRNA, or a genetic lesion in
a NOVX gene and to modulate NOVX activity. NOVX sequences are also useful
for identifying a cell or tissue type in a biological sample, to amplify
DNA sequences from very small biological samples such as tissues e.g.
hair or skin or body fluids in forensic biology and as primers and probes
for use in identifying and/or cloning NOVX homologues in other cell
types. The NOVX proteins are useful as an immunogen to generate
antibodies which are useful for diagnostically monitoring protein levels
and modulating NOVX activity. Cells comprising NOVX nucleic acids are
useful for producing non-human transgenic animals which are useful for
studying the function and/or activity of NOVX protein and for identifying
and/or evaluating modulators of NOVX protein activity. The NOVX nucleic
acids can be used in gene therapy. This polynucleotide sequence
represents a NOVX DNA sequence of the invention
Sequence 2463 BP; 691 A; 599 C; 580 G; 593 T; 0 U; 0 Other;
Query Match 41.4%; Score 412.2; DB 7; Length 2463;
Best Local Similarity 97.0%; Pred. No. 9.8e-82;
Matches 420; Conservative 0; Mismatches 13; Indels 0; Gaps 0;
QY 439 CTTTGTGTTTCCAGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGCAGA 498
DB 1998 CCGTAGTTTCAAGACACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGCAGA 2057

CC for treatment of muscle disease, e.g. Duchenne's muscular dystrophy
 CC (DMD). This sequence represents a mini-dystrophin sequence of the
 CC invention

XX Sequence 5339 BP; 1638 A; 1191 C; 1187 G; 1323 T; 0 U; 0 Other;

XX
 Query Match 41.2%; Score 410.8; DB 6; Length 5339;
 Best Local Similarity 98.3%; Pred. No. 2.4e-81;
 Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
 QY 450 AGGACACAATGAGGAAGTCTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 509
 DB 4491 AGGACACAATGAGGAAGTCTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 4550
 QY 510 CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTACAACCTCTGATT 569
 DB 4551 CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTACAACCTCTGATT 4610
 QY 570 CCGCATGGTTTTTATAATTTATATTCATACAAAGAGGATTTAGACAGTAAGAGTTTACAAG 629
 DB 4611 CCGCATGGTTTTTATAATTTATATTCATACAAAGAGGATTTAGACAGTAAGAGTTTACAAG 4670
 QY 630 AAATAAATCTATATTTTGTGAAGGAGTGTGTTTACTGATGATTTCAAGTATTTCT 689
 DB 4671 AAATAAATCTATATTTTGTGAAGGAGTGTGTTTACTGATGATTTCAAGTATTTCT 4730
 QY 690 AAGTCTGTTATTTGTTTAAATGTCAGATGGCAGGTTTTACACGTCCTATGCAATTTGACAAAA 749
 DB 4731 AAGTCTGTTATTTGTTTAAATGTCAGATGGCAGGTTTTACACGTCCTATGCAATTTGACAAAA 4790
 QY 750 AAGTTATAAGAAAACTACATGTAATAATCTTGATAGCTAAATAACTTGGCATTTCTTTATA 809
 DB 4791 AAGTTATAAGAAAACTACATGTAATAATCTTGATAGCTAAATAACTTGGCATTTCTTTATA 4850
 QY 810 TCGAACGCAATTTGGTGTGTTTAAAAATTTATAACAGTTATAAGAAAAAGATTTATAAAGG 869
 DB 4851 TCGAACGCAATTTGGTGTGTTTAAAAATTTATAACAGTTATAAGAAAAAGATTTATAAAGG 4910
 QY 870 AA 871
 DB 4911 AA 4912

RESULT 7
 ABK81997
 ID ABK81997 standard; DNA; 5417 BP.
 XX
 AC ABK81997;
 XX
 DT 13-AUG-2002 (first entry)
 XX
 DE DNA encoding mini-dystrophin protein deltaR4-R23.
 XX
 KW Mini-dystrophin peptide; spectrin-like repeat domain; muscle disease;
 KW Duchenne's muscular dystrophy; DMD; dystrophin; ds.
 XX
 OS Homo sapiens.
 OS Synthetic.
 XX
 PN WO200229056-A2.
 XX
 PD 11-APR-2002.
 XX
 PF 04-OCT-2001; 2001WO-US031126.
 XX
 PR 06-OCT-2000; 2000US-023848P.
 XX
 PA (UNMI) UNIV MICHIGAN.
 XX
 PI Chamberlain JS, Harper SQ;
 XX
 DR NPI; 2002-435334/46.
 XX

PT A composition for preparing therapeutic drugs, has a mini-dystrophin
 PT peptide comprising a specific number of spectrin-like repeat domains, or
 PT a nucleic acid sequence encoding the mini-dystrophin peptide.

XX Disclosure; Fig 12; 145pp; English.

XX
 The invention describes a composition comprising a mini-dystrophin
 CC peptide comprising a spectrin-like repeat domain, where the domain
 CC comprises n spectrin-like repeats, and contains no more than n spectrin-
 CC like repeats, where n is an even number between 4-24, or a nucleic acid
 CC encoding a mini-dystrophin peptide. The mini-dystrophin peptide or the
 CC polynucleotide encoding it is useful as a medicament, for preparing a
 CC drug for therapeutic application and in the preparation of a composition
 CC for treatment of muscle disease, e.g. Duchenne's muscular dystrophy
 CC (DMD). This sequence represents a mini-dystrophin sequence of the
 CC invention

XX Sequence 5417 BP; 1700 A; 1192 C; 1182 G; 1343 T; 0 U; 0 Other;

XX
 Query Match 41.2%; Score 410.8; DB 6; Length 5417;
 Best Local Similarity 98.3%; Pred. No. 2.4e-81;
 Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 450 AGGACACAATGAGGAAGTCTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 509
 DB 4569 AGGACACAATGAGGAAGTCTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 4628
 QY 510 CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTACAACCTCTGATT 569
 DB 4629 CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTACAACCTCTGATT 4688
 QY 570 CCGCATGGTTTTTATAATTTATATTCATACAAAGAGGATTTAGACAGTAAGAGTTTACAAG 629
 DB 4689 CCGCATGGTTTTTATAATTTATATTCATACAAAGAGGATTTAGACAGTAAGAGTTTACAAG 4748
 QY 630 AAATAAATCTATATTTTGTGAAGGAGTGTGTTTACTGATGATTTCAAGTATTTCT 689
 DB 4749 AAATAAATCTATATTTTGTGAAGGAGTGTGTTTACTGATGATTTCAAGTATTTCT 4808
 QY 690 AAGTCTGTTATTTGTTTAAATGTCAGATGGCAGGTTTTACACGTCCTATGCAATTTGACAAA 749
 DB 4809 AAGTCTGTTATTTGTTTAAATGTCAGATGGCAGGTTTTACACGTCCTATGCAATTTGACAAA 4868
 QY 750 AAGTTATAAGAAAACTACATGTAATAATCTTGATAGCTAAATAACTTGGCATTTCTTTATA 809
 DB 4869 AAGTTATAAGAAAACTACATGTAATAATCTTGATAGCTAAATAACTTGGCATTTCTTTATA 4928
 QY 810 TCGAACGCAATTTGGTGTGTTTAAAAATTTATAACAGTTATAAGAAAAAGATTTATAAAGG 869
 DB 4929 TCGAACGCAATTTGGTGTGTTTAAAAATTTATAACAGTTATAAGAAAAAGATTTATAAAGG 4988
 QY 870 AA 871
 DB 4989 AA 4990

RESULT 8
 ABK81999
 ID ABK81999 standard; DNA; 5462 BP.
 XX
 AC ABK81999;
 XX
 DT 13-AUG-2002 (first entry)
 XX
 DE DNA encoding mini-dystrophin protein deltaR2-R21+H3.
 XX
 KW Mini-dystrophin peptide; spectrin-like repeat domain; muscle disease;
 KW Duchenne's muscular dystrophy; DMD; dystrophin; ds.
 XX
 OS Homo sapiens.
 OS Synthetic.
 XX
 PN WO200229056-A2.

XX 11-APR-2002.
 XX 04-OCT-2001; 2001WO-US031126.
 XX 06-OCT-2000; 2000US-0238848P.
 XX (UNMI) UNIV MICHIGAN.
 XX Chamberlain JS, Harper SQ;
 XX WPI; 2002-435334/46.
 XX A composition for preparing therapeutic drugs, has a mini-dystrophin
 PT peptide comprising a specific number of spectrin-like repeat domains, or
 PT a nucleic acid sequence encoding the mini-dystrophin peptide.
 XX
 XX Disclosure; Fig 14; 145pp; English.
 XX
 XX The invention describes a composition comprising a mini-dystrophin
 CC peptide comprising a spectrin-like repeat domain, where the domain
 CC comprises n spectrin-like repeats, and contains no more than n spectrin-
 CC like repeats, where n is an even number between 4-24, or a nucleic acid
 CC encoding a mini-dystrophin peptide. The mini-dystrophin peptide or the
 CC polynucleotide encoding it is useful as a medicament, for preparing a
 CC drug for therapeutic application and in the preparation of a composition
 CC for treatment of muscle disease, e.g. Duchenne's muscular dystrophy
 CC (DMD). This sequence represents a mini-dystrophin sequence of the
 CC invention
 XX
 XX Sequence 5462 BP; 1668 A; 1225 C; 1212 G; 1357 T; 0 U; 0 Other;
 XX
 XX Query Match 41.2%; Score 410.8; DB 6; Length 5462;
 XX Best Local Similarity 98.3%; Pred. No. 2.4e-81;
 XX Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
 QY 450 AGGACACATGTAGGAAGCTCTTTCCACATGGCAGATGATTTGGCGAGCGATGGAGTC 509
 DB 4614 AGGACACATGTAGGAAGCTCTTTCCACATGGCAGATGATTTGGCGAGCGATGGAGTC 4673
 QY 510 CTTAGTATCAGTCATCAGATGAAGAGGAGCAGAAATTAATGTTTACAACTCCTGATT 569
 DB 4674 CTTAGTATCAGTCATCAGATGAAGAGGAGCAGAAATTAATGTTTACAACTCCTGATT 4733
 QY 570 CCGCATGGTTTTTAATTAATTCATCAACAAGAGGATTAGACAGTAAGATTTACAAAG 629
 DB 4734 CCGCATGGTTTTTAATTAATTCATCAACAAGAGGATTAGACAGTAAGATTTACAAAG 4793
 QY 630 AAATAAATCTATATTTTGTGAAGGCTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 689
 DB 4794 AAATAAATCTATATTTTGTGAAGGCTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 4853
 QY 690 AAGTCGTATTATTTTGTGAAGGCTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 749
 DB 4854 AAGTCGTATTATTTTGTGAAGGCTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 4913
 QY 750 AGCTTATAGAAACTACATGTAATCTTTGATAGCTAAATTAATCTTGCATTTCTTTATA 809
 DB 4914 AGCTTATAGAAACTACATGTAATCTTTGATAGCTAAATTAATCTTGCATTTCTTTATA 4973
 QY 810 TGAAGCGCATTTTGGGTTGTTTAAAAATTTTAAACAGTTTATAAAGAAAGATTTATAAAGG 869
 DB 4974 TGAAGCGCATTTTGGGTTGTTTAAAAATTTTAAACAGTTTATAAAGAAAGATTTATAAAGG 5033
 QY 870 AA 871
 DB 5034 AA 5035
 XX
 XX RESULT 9
 XX ABK82000
 XX ID ABK82000 standard; DNA; 8689 BP.
 XX

ABK82000;
 13-AUG-2002 (first entry)
 DNA encoding mini-dystrophin protein deltaH2-R19.
 Mini-dystrophin peptide; spectrin-like repeat domain; muscle disease;
 Duchenne's muscular dystrophy; DMD; dystrophin; ds.
 Homo sapiens.
 Synthetic.
 W0200229056-A2.
 11-APR-2002.
 04-OCT-2001; 2001WO-US031126.
 06-OCT-2000; 2000US-0238848P.
 (UNMI) UNIV MICHIGAN.
 Chamberlain JS, Harper SQ;
 WPI; 2002-435334/46.
 A composition for preparing therapeutic drugs, has a mini-dystrophin
 peptide comprising a specific number of spectrin-like repeat domains, or
 a nucleic acid sequence encoding the mini-dystrophin peptide.
 Disclosure; Fig 15; 145pp; English.
 The invention describes a composition comprising a mini-dystrophin
 peptide comprising a spectrin-like repeat domain, where the domain
 comprises n spectrin-like repeats, and contains no more than n spectrin-
 like repeats, where n is an even number between 4-24, or a nucleic acid
 encoding a mini-dystrophin peptide. The mini-dystrophin peptide or the
 polynucleotide encoding it is useful as a medicament, for preparing a
 drug for therapeutic application and in the preparation of a composition
 for treatment of muscle disease, e.g. Duchenne's muscular dystrophy
 (DMD). This sequence represents a mini-dystrophin sequence of the
 invention
 Sequence 8689 BP; 2721 A; 1804 C; 1861 G; 2303 T; 0 U; 0 Other;
 Query Match 41.2%; Score 410.8; DB 6; Length 8689;
 Best Local Similarity 98.3%; Pred. No. 2.7e-81;
 Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
 QY 450 AGGACACATGTAGGAAGCTCTTTCCACATGGCAGATGATTTGGCGAGCGATGGAGTC 509
 DB 5985 AGGACACATGTAGGAAGCTCTTTCCACATGGCAGATGATTTGGCGAGCGATGGAGTC 6044
 QY 510 CTTAGTATCAGTCATCAGATGAAGAGGAGCAGAAATTAATGTTTACAACTCCTGATT 569
 DB 6045 CTTAGTATCAGTCATCAGATGAAGAGGAGCAGAAATTAATGTTTACAACTCCTGATT 6104
 QY 570 CCGCATGGTTTTTAATTAATTCATCAACAAGAGGATTAGACAGTAAGATTTACAAAG 629
 DB 6105 CCGCATGGTTTTTAATTAATTCATCAACAAGAGGATTAGACAGTAAGATTTACAAAG 6164
 QY 630 AAATAAATCTATATTTTGTGAAGGCTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 689
 DB 6165 AAATAAATCTATATTTTGTGAAGGCTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 6224
 QY 690 AAGTCGTATTATTTTGTGAAGGCTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 749
 DB 6225 AAGTCGTATTATTTTGTGAAGGCTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 6284
 QY 750 AAGTTATAAGAAACTACATGTAATCTTTGATAGCTAAATTAATCTTGCATTTCTTTATA 809
 DB 6285 AAGTTATAAGAAACTACATGTAATCTTTGATAGCTAAATTAATCTTGCATTTCTTTATA 6344

Qy	630	AAATAAATCTATATTTTGTGGAAGGCTAGTGGTATTAATACTGTAGATTTTCAGTAGTTTCT	689
Db	8919	AAATAAATCTATATTTTGTGGAAGGCTAGTGGTATTAATACTGTAGATTTTCAGTAGTTTCT	8978
Qy	690	AACTCTGTTATTGTTTGTGTTAAACAATGGCAGGTTTACACGTCCTATGCAATTGTACAAAA	749
Db	8979	AACTCTGTTATTGTTTGTGTTAAACAATGGCAGGTTTACACGTCCTATGCAATTGTACAAAA	9038
Qy	750	AGTTTATAGAAACACATGTAATAATCTTGATAGCTAATAACTTGGCCATTTCTTTTATA	809
Db	9039	AGTTTATAGAAACACATGTAATAATCTTGATAGCTAATAACTTGGCCATTTCTTTTATA	9098
Qy	810	TGGAACGCAATTTTGGGTTGTTTAAAAATTTTAAACAGTTTATAAAGAAAGAAATTATAAAGG	869
Db	9099	TGGAACGCAATTTTGGGTTGTTTAAAAATTTTAAACAGTTTATAAAGAAAGAAATTATAAAGG	9158
Qy	870	AA 871	
Db	9159	AA 9160	
RESULT 11			
AAN90338			
ID	AAN90338 standard; cDNA; 12923 BP.		
XX	AAN90338;		
XX	29-MAR-1992 (first entry)		
XX	Sequence of human muscular dystrophy (MD) cDNA.		
XX	Dystrophin; muscular dystrophy; probe; antibody; diagnosis; prenatal;		
KW	heterozygote; gene therapy; genetic screening; foetal screening; ss.		
XX	Homo sapiens.		
XX	Key Location/Qualifiers		
PH	209..12923		
FT	/*tag= a		
XX	WO8906286-A.		
XX	13-JUL-1989.		
XX	16-DEC-1988; 88WO-US004504.		
XX	22-DEC-1987; 87US-00136618.		
XX	(CHIL-) CHILDRENS MED CENT.		
XX	Kunkel LM, Monaco A, Hoffman EP, Koenig M;		
XX	WPI; 1989-230587/30.		
DR	P-PSDB; AAF90373.		
XX	Muscular dystrophy gene - used for prepn. of probes, dystrophic		
FT	polypeptide and antibodies for diagnosis and therapy of muscular		
PT	dystrophy.		
XX	Disclosure; Fig 5; 68pp; English.		
XX	The inventors claim an MD probe comprising a purified ss NA SQ which		
CC	hybridises to at least a part of the MD gene; pure dystrophin (DS)		
CC	polypeptide, purified NA encoding DS and antibodies (Ab) to DS. The		
CC	probes are equal to or greater than 10b of one of 12 cDNA sequences		
CC	deposited as ATCC 58666-57677. The MD gene is human, or a murine Dmd gene		
XX	Sequence 12923 BP; 4296 A; 2613 C; 2963 G; 3044 T; 0 U; 7 Other;		
XX	Query Match 41.2%; Score 410.8; DB 1; Length 12923;		
XX	Best Local Similarity 98.3%; Pred. No. 2.9e-81;		
XX	Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;		

QY 450 AGGACACAAATGTAGGAAGTCTTTTCCACATGCGCAGATGATTTGGCAGAGCGATGGAGTC 509
 Db 11253 AGGACACAAATGTAGGAAGTCTTTTCCACATGCGCAGATGATTTGGCAGAGCGATGGAGTC 11312
 QY 510 CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATGAATGTTTCAACTCCTGATT 569
 Db 11313 CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATGAATGTTTCAACTCCTGATT 11372
 QY 570 CCGCATGCTTTTATAATATTATATACAAAGAGGATTTAGACAGTAAAGATTTCACAA 629
 Db 11373 CCGCATGCTTTTATAATATTATATACAAAGAGGATTTAGACAGTAAAGATTTCACAA 11432
 QY 630 AATAAATCTATATTTTGTGAAGGAGTGTGTTATATCTAGATTCAGTAGTTTCT 689
 Db 11433 AATAAATCTATATTTTGTGAAGGAGTGTGTTATATCTAGATTCAGTAGTTTCT 11492
 QY 690 AAGTCGTGTTTGTGTTTAACTGCGAGGTTTTCACAGTCTATGCAATTTGTACAAA 749
 Db 11493 AAGTCGTGTTTGTGTTTAACTGCGAGGTTTTCACAGTCTATGCAATTTGTACAAA 11552
 QY 750 AAGTTATAGAAACTACATGTAATCTTGTAGTAAATTAATCTGCAATTTCTTTATA 809
 Db 11553 AAGTTATAGAAACTACATGTAATCTTGTAGTAAATTAATCTGCAATTTCTTTATA 11612
 QY 810 TGGACGCAATTTGGGTGTTTAAATTTTAACTGTTTAAAGAAAGATTATAAAGG 869
 Db 11613 TGGACGCAATTTGGGTGTTTAAATTTTAACTGTTTAAAGAAAGATTATAAAGG 11672
 QY 870 AA 871
 Db 11673 AA 11674

RESULT 12
 ABR81959
 ID ABR81959 standard; DNA; 13957 BP.
 XX
 AC ABR81959;
 XX
 DT 13-AUG-2002 (first entry)
 XX
 DE cDNA encoding human dystrophin.
 XX
 KW Mini-dystrophin peptide; spectrin-like repeat domain; muscle disease;
 KW Duchenne's muscular dystrophy; DMD; dystrophin; human; gene; ds.
 XX
 OS Homo sapiens.
 XX
 PN WO200229056-A2.
 XX
 PD 11-APR-2002.
 XX
 XX 04-OCT-2001; 2001WO-US031126.
 XX
 XX 06-OCT-2000; 2000US-0238848P.
 XX
 XX (UNMI) UNIV MICHIGAN.
 XX
 XX Chamberlain JS, Harper SQ;
 XX
 XX WPI; 2002-435334/46.
 XX
 XX A composition for preparing therapeutic drugs, has a mini-dystrophin
 PT peptide comprising a specific number of spectrin-like repeat domains, or
 PT a nucleic acid sequence encoding the mini-dystrophin peptide.
 XX
 XX Example 2; Fig 1; 145pp; English.
 XX
 XX The invention describes a composition comprising a mini-dystrophin
 CC peptide comprising a spectrin-like repeat domain, where the domain
 CC comprises n spectrin-like repeats, and contains no more than n spectrin-
 CC like repeats, where n is an even number between 4-24, or a nucleic acid

CC encoding a mini-dystrophin peptide. The mini-dystrophin peptide or the
 CC polynucleotide encoding it is useful as a medicament, for preparing a
 CC drug for therapeutic application and in the preparation of a composition
 CC for treatment of muscle disease, e.g. Duchenne's muscular dystrophy
 CC (DMD). This sequence represents a human dystrophin polynucleotide
 CC sequence used in the creation of the mini-dystrophin peptides of the
 CC invention

SO Sequence 13957 BP; 4602 A; 2781 C; 3122 G; 3452 T; 0 U; 0 Other;

Query Match 41.2%; Score 410.8; DB 6; Length 13957;
 Best Local Similarity 98.3%; Pred. No. 3e-81;
 Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 450 AGGACACAAATGTAGGAAGTCTTTTCCACATGCGCAGATGATTTGGCAGAGCGATGGAGTC 509
 Db 11253 AGGACACAAATGTAGGAAGTCTTTTCCACATGCGCAGATGATTTGGCAGAGCGATGGAGTC 11312
 QY 510 CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATGAATGTTTCAACTCCTGATT 569
 Db 11313 CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATGAATGTTTCAACTCCTGATT 11372
 QY 570 CCGCATGCTTTTATAATATTATATACAAAGAGGATTTAGACAGTAAAGATTTCACAA 629
 Db 11373 CCGCATGCTTTTATAATATTATATACAAAGAGGATTTAGACAGTAAAGATTTCACAA 11432
 QY 630 AATAAATCTATATTTTGTGAAGGAGTGTGTTATATCTAGATTCAGTAGTTTCT 689
 Db 11433 AATAAATCTATATTTTGTGAAGGAGTGTGTTATATCTAGATTCAGTAGTTTCT 11492
 QY 690 AAGTCGTGTTTGTGTTTAACTGCGAGGTTTTCACAGTCTATGCAATTTGTACAAA 749
 Db 11493 AAGTCGTGTTTGTGTTTAACTGCGAGGTTTTCACAGTCTATGCAATTTGTACAAA 11552
 QY 750 AAGTTATAGAAACTACATGTAATCTTGTAGTAAATTAATCTGCAATTTCTTTATA 809
 Db 11553 AAGTTATAGAAACTACATGTAATCTTGTAGTAAATTAATCTGCAATTTCTTTATA 11612
 QY 810 TGGACGCAATTTGGGTGTTTAAATTTTAACTGTTTAAAGAAAGATTATAAAGG 869
 Db 11613 TGGACGCAATTTGGGTGTTTAAATTTTAACTGTTTAAAGAAAGATTATAAAGG 11672
 QY 870 AA 871
 Db 11673 AA 11674

RESULT 13
 ABR10904
 ID ABR10904 standard; cDNA; 13957 BP.
 XX
 AC ABR10904;
 XX
 DT 04-DEC-2002 (first entry)
 XX
 DE Human breast cancer associated coding sequence SEQ ID NO: 1038.
 XX
 KW Human; breast specific gene; breast cancer; differential expression;
 KW cytosstatic; gene therapy; Gene; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200259271-A2.
 XX
 PD 01-AUG-2002.
 XX
 XX 25-JAN-2002; 2002WO-US002176.
 XX
 XX 25-JAN-2001; 2001US-0263757P.
 PR 25-APR-2001; 2001US-0286090P.
 PR 23-MAY-2001; 2001US-0292517P.
 XX
 XX (GENE-) GENE LOGIC INC.

XX Orr MS, Nation M, Diggins JC, Zeng W;
XX WPI; 2002-674803/72.
XX Diagnosing breast cancer in a patient comprises detecting the level of
XX gene expression in cell or tissue samples, where a differential gene
XX expression is indicative of breast cancer.
XX
XX Claim 1; SEQ ID NO 1038; 260pp + Sequence Listing; English.
XX
XX The present invention relates to methods of diagnosing breast cancer in a
XX patient, which comprise detecting the level of expression in a tissue
XX sample of two or more genes selected from those shown in AB109867-
XX ABR1112, where a differential expression of the genes indicates breast
XX cancer. The methods are useful in diagnosing, treating, detecting the
XX progression, and in monitoring treatment of breast cancer in patients.
XX The methods are also useful as a screening tool for agents that modulate
XX the onset or progression of breast cancer. The breast cancer genes may be
XX used as diagnostic markers for the prediction or identification of the
XX malignant state of breast tissue, for confirming the type and progression
XX of cancer, and for drug screening and assays. The present sequence is a
XX coding sequence of the invention. Note: The sequence data for this patent
XX did not form part of the printed specification, but was obtained in
XX electronic format directly from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences
XX
XX Query Match 41.2%; Score 410.8; DB 6; Length 13957;
XX Best Local Similarity 98.3%; Pred. No. 3e-81;
XX Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
XX
XX QY 450 AGGACACATGTAGGAAGTCTTTCCACATGGCAGATGTTGGCAGAGCGATGGAGTC 509
XX DB 11253 AGGACACATGTAGGAAGTCTTTCCACATGGCAGATGTTGGCAGAGCGATGGAGTC 11312
XX
XX QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACCTCTGATT 569
XX DB 11313 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACCTCTGATT 11372
XX
XX QY 570 CCGCATGGTCTTTTATAATTTATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
XX DB 11373 CCGCATGGTCTTTTATAATTTATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG 11432
XX
XX QY 630 AAATAAATCTATATTTTGTGAAGGAGTGTGTTATATCTATCTAGATTTTCAGTAGTTTCT 689
XX DB 11433 AAATAAATCTATATTTTGTGAAGGAGTGTGTTATATCTATCTAGATTTTCAGTAGTTTCT 11492
XX
XX QY 690 AAGTCTGTTATTTGTTTAAACAATGGCAGGTTTACAGCTCTATGCAATTTGACAAAA 749
XX DB 11493 AAGTCTGTTATTTGTTTAAACAATGGCAGGTTTACAGCTCTATGCAATTTGACAAAA 11552
XX
XX QY 750 AAGTTTATAAGAAAACTACATGTAATAATCTTGATAGCTAAATTAACCTTGCATTTCTTTATA 809
XX DB 11553 AAGTTTATAAGAAAACTACATGTAATAATCTTGATAGCTAAATTAACCTTGCATTTCTTTATA 11612
XX
XX QY 810 TCGAAGCGAATTTGGTGTGTTTAAATAATTAACAGTTTAAAGAAATTAATAAGG 869
XX DB 11613 TCGAAGCGAATTTGGTGTGTTTAAATAATTAACAGTTTAAAGAAATTAATAAGG 11672
XX
XX QY 870 AA 871
XX DB 11673 AA 11674
XX
XX RESULT 14
XX ABN95786
XX ID ABN95786 standard; DNA; 13957 BP.
XX AC ABN95786;
XX DT 13-AUG-2002 (first entry)

XX Gene #2284 used to diagnose liver cancer.
XX DE
XX Gene: liver cancer; de; hepatocellular carcinoma; hepatotrophic;
XX metastatic liver tumour; cytostatic; expression profile; disease state;
XX disease progression; drug toxicity; drug efficacy; drug metabolism.
XX OS
XX Homo sapiens.
XX PN WO200229103-A2.
XX PD 11-APR-2002.
XX EF 02-OCT-2001; 2001WO-US030589.
XX FR 02-OCT-2000; 2000US-0237054P.
XX PA (GENE-) GENE LOGIC INC.
XX PI Horne D, Alvares C, Peres-Da-Silva S, Vockley JG;
XX WPI; 2002-426119/45.
XX DR
XX Diagnosing and detecting the progression of liver cancer, hepatocellular
XX carcinoma or metastatic liver tumor in a patient, involves detecting the
XX level of expression of two or more genes in a liver tissue sample.
XX
XX Claim 1; SEQ ID NO 2284; 298pp; English.
XX
XX The invention relates to a novel method for diagnosing and detecting the
XX progression of liver cancer, hepatocellular carcinoma or metastatic liver
XX tumour in a patient, and differentiating metastatic liver cancer from
XX hepatocellular carcinoma in a patient, involving detecting the level of
XX expression of two or more genes represented in ABN93503-ABN97455 in a
XX tissue sample. The method of the invention has hepatotrophic, and
XX cytostatic activity. The method is useful for diagnosing and detecting
XX the progression of liver cancer, hepatocellular carcinoma and metastatic
XX liver carcinoma in a patient. The method is useful for identifying
XX expression profiles which serve as useful diagnostic markers as well as
XX markers that can be used to monitor disease states, disease progression,
XX drug toxicity, drug efficacy and drug metabolism. Note: The sequence data
XX for this patent did not form part of the printed specification, but was
XX obtained in electronic format directly from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences
XX
XX Sequence 13957 BP; 4602 A; 2781 C; 3122 G; 3452 T; 0 U; 0 Other;
XX
XX Query Match 41.2%; Score 410.8; DB 6; Length 13957;
XX Best Local Similarity 98.3%; Pred. No. 3e-81;
XX Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
XX
XX QY 450 AGGACACATGTAGGAAGTCTTTCCACATGGCAGATGTTGGCAGAGCGATGGAGTC 509
XX DB 11253 AGGACACATGTAGGAAGTCTTTCCACATGGCAGATGTTGGCAGAGCGATGGAGTC 11312
XX
XX QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACCTCTGATT 569
XX DB 11313 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACCTCTGATT 11372
XX
XX QY 570 CCGCATGGTCTTTTATAATTTATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
XX DB 11373 CCGCATGGTCTTTTATAATTTATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG 11432
XX
XX QY 630 AAATAAATCTATATTTTGTGAAGGAGTGTGTTATATCTATCTAGATTTTCAGTAGTTTCT 689
XX DB 11433 AAATAAATCTATATTTTGTGAAGGAGTGTGTTATATCTATCTAGATTTTCAGTAGTTTCT 11492
XX
XX QY 690 AAGTCTGTTATTTGTTTAAACAATGGCAGGTTTACAGCTCTATGCAATTTGACAAAA 749
XX DB 11493 AAGTCTGTTATTTGTTTAAACAATGGCAGGTTTACAGCTCTATGCAATTTGACAAAA 11552
XX
XX QY 750 AAGTTTATAAGAAAACTACATGTAATAATCTTGATAGCTAAATTAACCTTGCATTTCTTTATA 809
XX DB 11553 AAGTTTATAAGAAAACTACATGTAATAATCTTGATAGCTAAATTAACCTTGCATTTCTTTATA 11612
XX
XX QY 810 TCGAAGCGAATTTGGTGTGTTTAAATAATTAACAGTTTAAAGAAATTAATAAGG 869
XX DB 11613 TCGAAGCGAATTTGGTGTGTTTAAATAATTAACAGTTTAAAGAAATTAATAAGG 11672
XX
XX QY 870 AA 871
XX DB 11673 AA 11674
XX
XX RESULT 14
XX ABN95786
XX ID ABN95786 standard; DNA; 13957 BP.
XX AC ABN95786;
XX DT 13-AUG-2002 (first entry)

Db 11553 AAGTTATAGAAAACACATGTAATCTTGATGCTAAATTAACCTGGCCATTTCTTTATA 11612
 QY 810 TCGAAGCGCATTTGGTCTTTTAAATTTATACAGTTATTAAGAAATTAAGG 869
 Db 11613 TCGAAGCGCATTTGGTCTTTTAAATTTATACAGTTATTAAGAAATTTGTAACCTA 11672
 QY 870 AA 871
 Db 11673 AA 11674
 RESULT 15
 ABS69900
 ID ABS69900 standard; DNA; 13957 BP.
 XX ABS69900;
 AC ABS69900;
 XX 21-NOV-2002 (first entry)
 XX Human dystrophin gene.
 DE
 XX Vector; adenovirus; adeno-associated; adenosine deaminase gene; receptor;
 KW adenosine deaminase deficiency; severe combined immune deficiency; PAH;
 KW beta-chain; haemoglobin gene; beta-thalassaemia; sickle cell disease;
 KW low density lipoprotein gene; familial hypercholesterolaemia;
 KW hypoxanthine-guanine phosphoribosyltransferase; Lesch-Nyhan syndrome;
 KW phenylalanine hydroxylase gene; gene therapy; phenylketonuria;
 KW dystrophin gene; muscular dystrophy; cystic fibrosis; immunostimulant;
 KW human cystic fibrosis transmembrane conductance regulator gene;
 KW antianemic; antilipemic; nootropic; cyostatic; dermatological; human;
 KW alpha-1-antitrypsin; lysosomal glucocerebrosidase; ADA; HPRT;
 KW lysosomal arylsulphatase A; ornithine transcarbamylase; ARSA; OTC; NP;
 KW purin nucleoside phosphorylase; gene; ds.
 XX Homo sapiens.
 OS
 XX US2002102731-A1.
 PN
 XX 01-AUG-2002.
 PD
 XX 12-FEB-2001; 2001US-00782378.
 PF
 XX 02-OCT-2000; 2000US-0237747P.
 PR
 XX (JYNY) UNIV NEW YORK STATE RES FOUND.
 PA
 XX Hearing P, Bahou WF, Sandalon Z, Gnatenko DV;
 XX WPI; 2002-690619/74.
 DR
 XX Producing vector, by introducing vector having nucleotide sequence,
 PT adenovirus inverted terminal repeats and packaging sequence, and adeno-
 PT associated virus terminal repeat, into cell, and culturing cell.
 XX
 PS Disclosure; Page 122-128; 191pp; English.
 XX
 CC The present invention relates to a new method of producing a vector. The
 CC method involves introducing recombinant vector having nucleotide sequence
 CC (NS) having 5' and 3' end, left and right inverted terminal repeats of
 CC adenovirus flanking NS, adenovirus packaging sequence linked to inverted
 CC terminal repeat, and adeno-associated virus terminal repeat linked to 3'
 CC end of NS, into cell expressing adenovirus early gene lacking from vector
 CC ; and culturing cell to produce another vector. The method is useful for
 CC generating vectors, especially mad vectors. The method is useful in
 CC transferring nucleotide sequences of interest into a cell, for gene
 CC transfer applications (e.g. gene therapy) in vitro, ex vivo and in vivo.
 CC The nucleotide sequences are useful for treating diseases associated with
 CC it, i.e. adenosine deaminase gene associated with adenosine deaminase
 CC deficiency with severe combined immune deficiency, beta-chain of
 CC haemoglobin gene associated with beta-thalassaemia and sickle cell
 CC disease, receptor for low density lipoprotein gene associated with
 CC familial hypercholesterolaemia, hypoxanthine-guanine
 CC phosphoribosyltransferase associated with Lesch-Nyhan syndrome,

CC phenylalanine hydroxylase (PAH) gene associated with phenylketonuria;
 CC dystrophin gene associated with muscular dystrophy, and human cystic
 CC fibrosis transmembrane conductance regulator gene associated with cystic
 CC fibrosis. The present nucleic acid sequence represents a human disease
 CC gene sequence that was used in the methods of the invention
 XX
 SQ Sequence 13957 BP; 4602 A; 2781 C; 3122 G; 3452 T; 0 U; 0 Other;
 Query Match 41.2%; Score 410.8; DB 6; Length 13957;
 Best Local Similarity 98.3%; Pred. NO. 3e-81; 7; Indels 0; Gaps 0;
 Matches 415; Conservative 0; Mismatches 0;
 QY 450 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGGCGAGCGATGGAGTC 509
 Db 11253 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGGCGAGCGATGGAGTC 11312
 QY 510 CTTAGTATCATGATGACAGATGAAGAGGAGCAGATGAATGTTTACAACTCCTGATT 569
 Db 11313 CTTAGTATCATGATGACAGATGAAGAGGAGCAGATGAATGTTTACAACTCCTGATT 11372
 QY 570 CCCGCATGGTCTTTTATAATATTCATACACAAAGAGGATTTAGACAGTAAAGTTTCAAG 629
 Db 11373 CCCGCATGGTCTTTTATAATATTCATACACAAAGAGGATTTAGACAGTAAAGTTTCAAG 11432
 QY 630 AATATAATCTATATTTTGTGAAGGTAGTGGTATTATCTAGTATTTTACAGTATTCT 689
 Db 11433 AATATAATCTATATTTTGTGAAGGTAGTGGTATTATCTAGTATTTTACAGTATTCT 11492
 QY 690 AAGTCTCTTATTTGTTTAACTAATGGCAGGTTTACACGCTCTATGCAATTGTCAAAA 749
 Db 11493 AAGTCTCTTATTTGTTTAACTAATGGCAGGTTTACACGCTCTATGCAATTGTCAAAA 11552
 QY 750 AAGTTATAAGAAAACACTACATGTAATCTTTGATAGCTAATAAATCTTGCCATTTCTTTATA 809
 Db 11553 AAGTTATAAGAAAACACTACATGTAATCTTTGATAGCTAATAAATCTTGCCATTTCTTTATA 11612
 QY 810 TGGACCGCATTTGGGTTGTTTAAATTTTATAACAGTTTATAAGAAAGAAATTTATAAAGG 869
 Db 11613 TGGACCGCATTTGGGTTGTTTAAATTTTATAACAGTTTATAAGAAAGAAATTTATAAAGG 11672
 QY 870 AA 871
 Db 11673 AA 11674

Search completed: April 6, 2004, 11:42:39
 Job time : 473.188 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 08:36:01 ; Search time 110.764 Seconds
(without alignments)
4990.154 Million cell updates/sec

Title: US-09-966-264D-2
Perfect score: 996
Sequence: 1 ggggttgatgatagtaaa.....gtgttgatgtaattaatt 996

Scoring table: IDENTITY_NUC

Gapop 10.0 , Capext 1.0

Searched: 682709 seqs, 277475446 residues

Total number of hits satisfying chosen parameters: 1365418

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents NA:*
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3: /cgn2_6/ptodata/2/ina/6A COMB.seq:*
4: /cgn2_6/ptodata/2/ina/6B COMB.seq:*
5: /cgn2_6/ptodata/2/ina/PTUS COMB.seq:*
6: /cgn2_6/ptodata/2/ina/backfiles1.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	409.2	41.1	13977	4	US-09-484-970B-60
C 2	374.2	37.6	19307	3	US-08-836-022A-10
C 3	374.2	37.6	19307	3	US-09-427-048A-10
C 4	140.4	14.1	13977	4	US-09-484-970B-60
5	101.4	10.2	19307	3	US-08-836-022A-10
6	101.4	10.2	19307	3	US-09-427-048A-10
C 7	45	4.5	615	3	US-08-998-416-186
8	44	4.4	8093	4	US-10-204-708-31
C 9	43.5	4.4	832	4	US-09-621-976-2813
10	43	4.3	1830121	4	US-09-557-884-1
11	43	4.3	1830121	4	US-09-643-990A-1
C 12	42.8	4.3	7218	1	US-08-232-463-14
13	42.4	4.3	19124	2	US-08-487-826B-13
14	42	4.2	5156	2	US-09-091-432-3
15	42	4.2	5156	4	US-09-387-663-3
16	42	4.2	5156	4	US-09-214-139B-3
C 17	41.5	4.2	640681	4	US-09-790-988-1
18	41.2	4.1	11049	4	US-10-204-708-21
19	41	4.1	15567	4	US-09-627-376-3
20	40.6	4.1	615	3	US-08-998-416-186
21	40.6	4.1	636	3	US-08-998-416-1137
22	40.6	4.1	837	3	US-08-998-416-288
23	40.4	4.1	5152	4	US-10-204-708-73
C 24	40.4	4.1	6124	4	US-08-213-419B-3
C 25	40.4	4.1	319608	4	US-09-539-333B-1
C 26	40.4	4.1	319608	4	US-09-679-409-1
27	40	4.0	6124	4	US-08-213-419B-3

ALIGNMENTS

RESULT 1

US-09-484-970B-60
; Sequence 60, Application US/09484970B
; Patent No. 6426186
; GENERAL INFORMATION:
; APPLICANT: Jones, Karen A.
; APPLICANT: Volkmutch, Wayne
; APPLICANT: Walker, Michael G.
; TITLE OF INVENTION: BONE REMODELING GENES
; FILE REFERENCE: PB-0014 US
; CURRENT APPLICATION NUMBER: US/09/484,970B
; CURRENT FILING DATE: 2000-01-18
; NUMBER OF SEQ ID NOS: 172
; SOFTWARE: PERL Program
; SEQ ID NO 60
; LENGTH: 13977
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: Incyte ID No. 6426186 229357.11CBI
; NAME/KEY: unsure
; LOCATION: 11721-11761, 12294, 13969
; OTHER INFORMATION: a, t, c, g, or other
US-09-484-970B-60

Query Match Similarity 41.1%; Score 409.2; DB 4; Length 13977;
Best Local Similarity 98.1%; Pred. No. 1.6e-92; Indels 0; Gaps 0;
Matches 414; Conservative 0; Mismatches 8

QY	450	AGGACACAATGTAGGAAGTCTTTCCACATGGCAGATGATTGGCGAGCGATGGAGTC	509
DB	11254	AGGACACAATGTAGGAAGTCTTTCCACATGGCAGATGATTGGCGAGCGATGGAGTC	11313
QY	510	CTTAGTATCATGATGACAGATGAAGAGGAGCAGATTAATGTTTACAACTCCTGATT	569
DB	11314	CTTAGTATCATGATGACAGATGAAGAGGAGCAGATTAATGTTTACAACTCCTGATT	11373
QY	570	CCCGCATCGTTTTTATATATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG	629
DB	11374	CCCGCATCGTTTTTATATATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG	11433
QY	630	AAATAAATCTATATTTTGTGAAGGAGTAGTGATTAATATCTGTAGATTTCAAGTAGTTCT	689
DB	11434	AAATAAATCTATATTTTGTGAAGGAGTAGTGATTAATATCTGTAGATTTCAAGTAGTTCT	11493
QY	690	AAGTCGTATTGTTTGTAAACAATGCGAGTTTACACGCTTATGCAATTTGTACAAA	749
DB	11494	AAGTCGTATTGTTTGTAAACAATGCGAGTTTACACGCTTATGCAATTTGTACAAA	11553

Sequence 7, Appli
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Sequence 1, Appli
Sequence 5, Appli
Sequence 23, Appli
Sequence 82, Appli
Sequence 46, Appli
Sequence 1, Appli
Sequence 1, Appli
Sequence 2813, Ap
Sequence 1, Appli
Patent No. 5256558
Sequence 6, Appli
Sequence 3, Appli
Sequence 58, Appli
Sequence 1797, Ap
Sequence 1797, Ap
Sequence 1797, Ap

QY	750	AGTTATATAGAAAACTACATGCTGTAATATCTTGATAGCTAAATTAACCTTGCATTTCTTTATA	809
DB	11554	AGTTATATAGAAAACTACATGCTGTAATATCTTGATAGCTAAATTAACCTTGCATTTCTTTATA	11613
QY	810	TGGAACGCCATTTTGGGTTGTTTAAAAATTTATAACAGTTTATAAAGAAAGAAATTTATAAAGG	869
DB	11614	TGGAACGCCATTTTGGGTTGTTTAAAAATTTATAACAGTTTATAAAGAAAGAAATTTATAAAGG	11673
QY	870	AA 871	
DB	11674	AA 11675	
RESULT 2			
US-08-836-022A-10/c			
; Sequence 10, Application US/08836022A			
; Patent No. 6001557			
; GENERAL INFORMATION:			
; APPLICANT: Trustees of the University of Pennsylvania			
; APPLICANT: Wilson, James M.			
; APPLICANT: Fisher, Krishna J.			
; APPLICANT: Chen, Shu-Jen			
; APPLICANT: Weitzman, Matthew			
; TITLE OF INVENTION: Improved Adenovirus Virus and			
; NUMBER OF SEQUENCES: 10			
; CORRESPONDENCE ADDRESS:			
; ADDRESSEE: Howson and Howson			
; STREET: Spring House Corporate Cntr, P O Box 457			
; CITY: Spring House			
; STATE: Pennsylvania			
; COUNTRY: USA			
; ZIP: 19477			
; COMPUTER READABLE FORM:			
; MEDIUM TYPE: Floppy disk			
; COMPUTER: IBM PC compatible			
; OPERATING SYSTEM: PC-DOS/MS-DOS			
; SOFTWARE: Patent In Release #1.0, Version #1.30			
; CURRENT APPLICATION DATA:			
; APPLICATION NUMBER: US/08/836,022A			
; FILING DATE:			
; CLASSIFICATION: 435			
; PRIOR APPLICATION DATA:			
; APPLICATION NUMBER: US 08/331,381			
; FILING DATE: 28-OCT-1994			
; ATTORNEY/AGENT INFORMATION:			
; NAME: Bak, Mary E.			
; REGISTRATION NUMBER: 31,215			
; REFERENCE/DOCKET NUMBER: GNPVN.008PCT			
; TELECOMMUNICATION INFORMATION:			
; TELEPHONE: 215-540-9200			
; TELEFAX: 215-540-5818			
; INFORMATION FOR SEQ ID NO: 10:			
; SEQUENCE CHARACTERISTICS:			
; LENGTH: 19307 base pairs			
; TYPE: nucleic acid			
; STRANDEDNESS: double			
; TOPOLOGY: unknown			
; MOLECULE TYPE: CDNA			
US-08-836-022A-10			
Query Match 37.6%; Score 374.2; DB 3; Length 19307;			
Best Local Similarity 96.0%; Pred No. 9.5e-84;			
Matches 406; Conservative 0; Mismatches 13; Indels 4; Gaps 2;			
QY	450	AGGCACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGCGAGCGCATGGAGTC	509
DB	3441	AGGCACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGCGAGCGCATGGAGTC	3382
QY	510	CTTAGTATCACTCATGACAGATGAGAGGAGCAGAGATAAATGTTTACCACTCTTGATT	569
DB	3381	CTTAGTATCACTCATGACAGATGAGAGGAGCAGAGATAAATGTTTACCACTCTTGATT	3322
QY	570	CCCCGATGTTTATTAATATTCATACAAACAAGAGGATTAGACAGTAACAGTTTACAAG	629

Db	3321	CCGCAATGGTTTATTAATATTCGTACACAAAGGATTACAGCTAAGAGTTTACAAG	3362
Qy	630	AAAT-AAATCTATATTTTGTGAAGGGTAGTGGTATATATCTGTAGATTTCAGTAGTTTC	688
Db	3261	AAATAAAATCTATATTTTGTGAAGGGTAGTGGTACTATCTGTAGATTTCAGTAGTTTC	3202
Qy	689	TAAGTCGTATTGTTTGTTAACAATGCGAGGTTTACACGCTCTATGCAATTGTACAAA	748
Db	3201	TAAGTCGTATTGTTTGTTAACAATGCGAGGTTTACACGCTCTATGCAATTGTACAAA	3142
Qy	749	AAAGTTTATAAGAAAACATCATGTAATCTTGATAGCTAAATAACCTGCGATTTCTTTAT	808
Db	3141	AAAGTTTAAAAAGAAA--ACATGTAAAAATCTTGATAGCTAAATAACCTGCGATTTCTTTAT	3085
Qy	809	ATGGAACGCAATTTGGGTGTTTAAAAATTTATAACAGTTTATAAAGAAAGAAATATAAAG	868
Db	3084	ATGGAACGCAATTTGGGTGTTTAAAAATTTATAACAGTTTATAAAGAAAGAAATATAAAG	3025
Qy	869	GAA 871	
Db	3024	AAA 3022	

RESULT 3
 US-09-427-048A-10/c
 ; Sequence 10, Application US/09427048A
 ; Patent No. 6203975
 ; GENERAL INFORMATION:
 ; APPLICANT: Trustees of the University of Pennsylvania
 ; Wilson, James M.
 ; Fisher, Krishna J.
 ; Chen, Shu-Jen
 ; Weitzman, Matthew
 ; TITLE OF INVENTION: Improved Adenovirus Virus and
 ; Methods of Use Thereof
 ; NUMBER OF SEQUENCES: 10
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Howson and Howson
 ; STREET: Spring House Corporate Cntr, P O Box 457
 ; CITY: Spring House
 ; STATE: Pennsylvania
 ; COUNTRY: USA
 ; ZIP: 19477
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: Patent in Release #1.0, Version #1.30
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/09/427,048A
 ; FILING DATE: 21-Oct-1999
 ; CLASSIFICATION: <Unknown>
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: 08/836,022
 ; FILING DATE: <Unknown>
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Bak, Mary E.
 ; REGISTRATION NUMBER: 31,215
 ; REFERENCE/DOCKET NUMBER: GNPVN.008PCT
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: 215-540-3200
 ; TELEFAX: 215-540-5818
 ; INFORMATION FOR SEQ ID NO: 10:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 19307 base pairs
 ; TYPE: nucleic acid
 ; STRANDEDNESS: double
 ; TOPOLOGY: unknown
 ; MOLECULE TYPE: cDNA
 ; SEQUENCE DESCRIPTION: SEQ ID NO: 10:
 US-09-427-048A-10

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SEQUENCE DESCRIPTION: SEQ ID NO: 10
US-09-427-048A-10

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Query Match      37.6%; Score 374.2; DB 3; Length 19307;
Best Local Similarity 96.0%; Pred. No. 9.5e-84;
Matches 406; Conservative 0; Mismatches 13; Indels 4; Gaps 2;

QY 450 AGGACACATGATGAGGAGTCTTTTCCACATGCGCAGATGTTGGCAGAGCGATGGAGTC 509
DB 3441 AGGACACATGATGAGGAGTCTTTTCCACATGCGCAGATGTTGGCAGAGCGATGGAGTC 3382

QY 510 CTTAGTATCAGTCATGACAGATGAAAGAGGAGCGAGATTAATGTTTACAACTCTCTGATT 569
DB 3381 CTTAGTATCAGTCATGACAGATGAAAGAGGAGCGAGATTAATGTTTACAACTCTCTGATT 3322

QY 570 CCCGATGTTTTTATAATATTATACACAAAGAGGATTAGACAGTAAGAGTTTACAAAG 629
DB 3321 CCCGATGTTTTTATAATATTATCGTACACAAAGAGGATTAGACAGTAAGAGTTTACAAAG 3262

QY 630 AAAT-AAATCTATATTTTGTGAAGGAGTGGTATTAATCTAGTATGATTTTCAAGTATTC 688
DB 3261 AAATAAATCTATATTTTGTGAAGGAGTGGTATTAATCTAGTATGATTTTCAAGTATTC 3202

QY 689 TAAGTCTGTATGTTTGTAAACATGCGAGGTTTACAGCTCTATGCAATTTACAAA 748
DB 3201 TAAGTCTGTATGTTTGTAAACATGCGAGGTTTACAGCTCTATGCAATTTACAAA 3142

QY 749 AAAGTTATAAGAAAACATCATGTAATAATCTTCATAGCTAAATAACTTCCCATTTCTTTAT 808
DB 3141 AAAGTTATAAGAAAACATCATGTAATAATCTTCATAGCTAAATAACTTCCCATTTCTTTAT 3085

QY 809 ATGGAACGATTTTGGTGTGTTTAAATTTTAAACATGTTTAAAGAGGATTAATAAG 868
DB 3084 ATGGAACGATTTTGGTGTGTTTAAATTTTAAACATGTTTAAAGAGGATTTAAACT 3025

QY 869 GAA 871
DB 3024 AAA 3022

RESULT 4
US-09-484-970B-60/c
; Sequence 60, Application US/09484970B
; Patent No. 6426186
; GENERAL INFORMATION:
; APPLICANT: Jones, Karen A.
; APPLICANT: Volkmut, Wayne
; APPLICANT: Walker, Michael G.
; TITLE OF INVENTION: BONE REMODELING GENES
; FILE REFERENCE: PB-0014 US
; CURRENT APPLICATION NUMBER: US/09/484,970B
; CURRENT FILING DATE: 2000-01-18
; NUMBER OF SEQ ID NOS: 172
; SOFTWARE: PERL Program
; SEQ ID NO 60
; LENGTH: 13977
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; OTHER INFORMATION: Incyte ID No. 6426186 229357.11CB1
; NAME/KEY: unsure
; LOCATION: 11721-11761, 12294, 13969
; OTHER INFORMATION: a, t, c, g, or other
US-09-484-970B-60

Query Match      14.1%; Score 140.4; DB 4; Length 13977;
Best Local Similarity 90.4%; Pred. No. 1.8e-25;
Matches 161; Conservative 0; Mismatches 16; Indels 1; Gaps 1;

QY 820 TTGGGTGTTTAAATTTATACAGTTTATTAAGAAAGAAATTTAAAGGAAAAAGAAAA 879
DB 13346 TTGGGTGTTTCTTTGAAATTTATGAGGAAAGAAAGAAATTTAAAGGAAAAAGAAAA 13287

QY 880 TAACGAATGGACAGTGGTGAAGCTGTGAATCTAGGTGTGCACATTTATCAGGAACACC 939

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DB 13286 TAACGCAATGCAAGTGGTGAAGCTGTGAATCTAGGTGTGCACATTTATCAGGAACACC 13227
QY 940 CCAAAACCAAGTGAAGTGAATAATAGCATGAGAAG-CCGTGTTTGTATGTTAATTAATT 996
DB 13226 CCAAAACCAAGTGAAGTGAATAATAGCATGAGAAGCCCGTGTGATGTTAATTAATT 13169

RESULT 5
US-08-836-022A-10
; Sequence 10, Application US/08836022A
; Patent No. 6001557
; GENERAL INFORMATION:
; APPLICANT: Trustees of the University of Pennsylvania
; APPLICANT: Wilson, James M.
; APPLICANT: Fisher, Krishna J.
; APPLICANT: Chen, Shu-Jen
; APPLICANT: Weitzman, Matthew
; TITLE OF INVENTION: Improved Adenovirus Virus and
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Howson and Howson
; STREET: Spring House Corporate Cntr, P O Box 457
; CITY: Spring House
; STATE: Pennsylvania
; COUNTRY: USA
; ZIP: 19477
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/836,022A
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/331,381
; FILING DATE: 28-OCT-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Bak, Mary E.
; REGISTRATION NUMBER: 31,215
; REFERENCE/DOCKET NUMBER: GNVN.008PCT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 215-540-9200
; TELEFAX: 215-540-5818
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 19307 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: unknown
; MOLECULE TYPE: cDNA
US-08-836-022A-10

Query Match      10.2%; Score 101.4; DB 3; Length 19307;
Best Local Similarity 81.8%; Pred. No. 1e-15;
Matches 117; Conservative 0; Mismatches 26; Indels 0; Gaps 0;

QY 851 AAGAAAGAAATTTAAAGGAAAAAGAAATAACGCAATGGAACAAGTGGTGAAGCTGTGAA 910
DB 1480 AATTATAATGAAAAAGAAACTGGTGTCCACACACAGCAAGTGGTGAAGTGTGAA 1539

QY 911 CTCAGGTGTGCACATTTATCAGGAACACCCCAAAACCAAGTGAAGTGAAGTGAAGTGA 970
DB 1540 ATTAGGTGTGCACATTTATCAGGAACACCCCAAAACCAAGTGAAGTGAAGTGAAGTGA 1599

QY 971 GAAGCCGTGTTTGTATGTTAATTA 993
DB 1600 GAAGCTGTGTTTGTATGTTAATTA 1622

RESULT 6
US-09-427-048A-10

```

APPLICANT: Wendland, Jürgen
APPLICANT: Knechtle, Philipp
APPLICANT: Reibschung, Corinne
TITLE OF INVENTION: GENOMIC DNA SEQUENCES OF ASHBYA GOSYPII
TITLE OF INVENTION: AND USES THEREOF
NUMBER OF SEQUENCES: 1152
CORRESPONDENCE ADDRESS:
ADDRESS: No. 6239264artis Corporation
STREET: 3054 Cornwallis Road
CITY: Research Triangle Park
STATE: No. 6239264th Carolina
COUNTRY: USA
ZIP: 27709

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/998,416
FILING DATE: 24-DEC-1997
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: CH 0016/97
FILING DATE: 31-DEC-1996
ATTORNEY/AGENT INFORMATION:
NAME: Weigs, J. Timothy
REGISTRATION NUMBER: 38,241
REFERENCE/DOCKET NUMBER: PF/5-30306/A/OGC1976
TELECOMMUNICATION INFORMATION:
TELEPHONE: 919-541-8587
TELEFAX: 919-541-8689

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; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; ORIGINAL SOURCE:
; ORGANISM: PAG1074RP
; US-08-998-416-186

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Query Match 4.5%; Score 45; DB 3; Length 615;
Best Local Similarity 48.0%; Pred. No. 0.037;
Matches 129; Conservative 0; Mismatches 140; Indels

593	QY	TATATATTCATACACAAAGAGGATTAGACAGCTAAGAGTTTACAGAAATAAATCTATA	642
576	DB	TAAATAATTAATTTAAATAAATAATTTAAATTAATAATAATAATAAGTGAATATTAAT	517
643	QY	TTTTTGTGAAGGGTAGTGGTATTATATCTGAGATTTTCAGTAGTTTCTAAGTCTGTTATTG	702
516	DB	TTATTAAATAATAATAAAAAATTAAATAAGAAATTAAGTTTAAAAATTATTTTAAATAA	457
703	QY	TTTTTCTTAAACAATGGCAGGTTTTACAGCTCTAGCAATTGTACAAAAAAGTTTATAAGAA	762
456	DB	ATTCCTTATAAAAAGTTTAAATAATAATAATAAACCAATATATTTATAAAAATAGTATTATA	397
763	QY	ACTACATGTAAAAATCTTGATAGCTAAATAACITGCCATTTCTTTATATCGAAGCGCATTTT	822
306	DB	ATATAAATAAACAATTTTATCAATATTTAAATAAATAATTTTAAATCTTTTAAATAAATAATTT	337

RESULT 8
US-10-204-708-31
; Sequence 31, Application US/10204708
; Patent No. 667731
; GENERAL INFORMATION:

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; APPLICANT: OLEK, Alexander
; APPLICANT: PIERBROCK, Christian
; APPLICANT: BERLIN, Kurt
; TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication
; TITLE OF INVENTION: by Assessing DNA Methylation
; FILE REFERENCE: 5013.1012
; CURRENT APPLICATION NUMBER: US/10/204,708
; CURRENT FILING DATE: 2003-05-06
; PRIOR APPLICATION NUMBER: PCT/EP01/03971
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: DE 10019058.8
; PRIOR FILING DATE: 2000-04-06
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; PRIOR APPLICATION NUMBER: DE 10032529.7
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: DE 10043826.1
; PRIOR FILING DATE: 2000-09-01
; NUMBER OF SEQ ID NOS: 98
; SEQ ID NO 31
; LENGTH: 8093
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-204-708-31

Query Match
Best Local Similarity 4.4%; Score 44; DB 4; Length 8093;
Matches 149; Conservative 0; Mismatches 155; Indels 2; Gaps 1;

QY 579 TTTTATAATATTCATCAACAAGAGGATTAGACAGTAAGAGTTTACAAGAAATAATC 638
DB 336 TTTTAAATATATTTATTGTTATATAGGATTAGTAATATATATTTTAAAAAATAA 395

QY 639 TATATTTTGTGAAGGTAGTGGTATATATCTAGATTTTCAAGTCTGTT 698
DB 396 AAGATGTTATGTTATAGTAATATTTAATATAGAAATAGTAAATATATTTT 455

QY 699 ATTGTTTGTAAACATGGCAGGTTTACACGCTCTATGCAATTCACAAAAAGTTATA 758
DB 456 TTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTT 515

QY 759 GAAAC--TACATGPAATCTGTAGTCTAAATCTGCAATTTCTTTATATGGAACG 816
DB 516 TTAAGGGTAAATTTAAATTTTGGTTATATTTTATTTTATTTTATTTTATTTATG 575

QY 817 CATTTTGGTGTGTTAAAAATTTATACAGTTTATAAGAAAGAAATTTATAAGGAAAGA 876
DB 576 TAAATTAATTTTATTAATTTTATAGTAAATATATTTGGAATTTTATTAATATGTAAGA 635

QY 877 AAATAA 882
DB 636 TTTTAA 641

RESULT 9
US-09-621-976-2813/c
; Sequence 2813, Application US/09621976
; Patent No. 6639063
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Jobert, S.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: ESTs and Encoded Human Proteins.
; FILE REFERENCE: GENSET.054PR2
; CURRENT APPLICATION NUMBER: US/09/621,976
; CURRENT FILING DATE: 2000-07-21
; NUMBER OF SEQ ID NOS: 19335
; SOFTWARE: Patent.pm
; SEQ ID NO 2813
; LENGTH: 832
; TYPE: DNA

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; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 235..399
US-09-621-976-2813

Query Match
Best Local Similarity 4.4%; Score 43.6; DB 4; Length 832;
Matches 67; Conservative 165; Mismatches 184; Indels 2; Gaps 1;

QY 576 TGTGTTTATAATATTCATCAACAAGAGGATTAGACAGTAAGAGTTTACAAGAAATAA 635
DB 433 TGACTTTGACTATGAACATACTCCAGATTGCTAAAGTACAAGAATTGTACCATATAA 374

QY 636 ATCTATATTTTGTGAAGGTAGTGGTATATATCTAGTATTCAGTATTCAGTAGTCT 695
DB 373 TTAATATTTTGTGTTTWWKWTWYTTTWTMMWKKARRRYYWKKSYVACASRYKKYTW 314

QY 696 G--TTATTTGTTTAAACAATGGCAGGTTTACACGCTCTATGCAATTTGACAAAAAAGT 753
DB 313 GWWYMWKEMSTRWYCYWCKCKVGRRCAYTWMARGMWSYAWGKWSMSMCT 254

QY 754 TATAAGAAAACTACATGTAATAATCTTATAGTCTAAATACTTGCCATTTCTTTATATGA 813
DB 253 RMYKKGSTYWTMCKTCAWYWKYKRMWSKTCWSGSGGYMTSYTSTRSYMYWAS 194

QY 814 ACGCATTTTGGTGTGTTTAAAAATTTATAACAGTTTATAAGAAAGAAATTTATAAGAAAA 873
DB 193 WMYTMCWNGRWSTYWMWGWKKWRYATTWRRAMWWAAATWYMWYMWKSSRGA 134

QY 874 AGAAATAACGCAATGGAACAAGTGTGAAGCTGTGAACCTCAGGTGTGCACAATTATCAGG 933
DB 133 AMYRRTMMWGVYVWKKSYRRTRCWAYAWKTRSYVWCKWCKRCKMMMMMAAYG 74

QY 934 AACACCCCAACCAAGTGAAGTGAATAATACATCAGAACGCGTGTGTTGATTAAT 991
DB 73 KTMGRACWKTFRWAWAMMMWMTMMMTTYTWRAMKRRWMMKRSNWMMAW 16

RESULT 10
US-557-884-1
; Sequence 1, Application US/09557884
; Patent No. 6506581
; GENERAL INFORMATION:
; APPLICANT: Fleischmann et al.
; TITLE OF INVENTION: The Nucleotide sequence of
; the Haemophilus influenzae Rd Genome, Fragments
; Thereof, and Uses Thereof
; NUMBER OF SEQUENCES: 1
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: MD
; COUNTRY: USA
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3 1/2 inch diskette
; COMPUTER: Dell Pentium
; OPERATING SYSTEM: MS DOS v6.22
; SOFTWARE: ASCII Text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/557,884
; FILING DATE: 25-Apr-2000
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/476,102
; FILING DATE: JUN-5-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Michelle S. Marks
; REGISTRATION NUMBER: 41,971
; REFERENCE/DOCKET NUMBER: PE186P3
; TELECOMMUNICATION INFORMATION:

```

```

; TELEPHONE: 301-309-8504
; TELEFAX: 301-309-8439
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1830121 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-557-884-1

Query Match          4.3%; Score 43; DB 4; Length 1830121;
Best Local Similarity 48.7%; Pred. No. 1.9;
Matches 115; Conservative 0; Mismatches 121; Indels 0; Gaps 0;

QY 738 AATTGTACAAAAAGTTTAAAGAAACTACATGTAAATCTTTGATAGTAAATAACTTGC 797
Db 413354 AAGTGTATACCAATATATAAGAGAGATGGAATAGTAAGATAGATAAATGCTGTT 413413

QY 798 CATTTCTTTATATGGAACGCAATTTTGGGTTGTTTAAAAATTTTAAACAGTTTATAAGAAA 857
Db 413414 TAAAGTTTAAATTTGATTTTCTATAGGTTGCTCAATTAATTAATAACTTTGTGATAGAAA 413473

QY 858 GAATTATAAGAAAAAGAAATACGCAATGGAAGTGTGAGTGAATAGCATGAGAA 917
Db 413474 AAATAATAGTTAATAATAACAACAATTAACATGTAATGCTTGAATCTGCTGCGCAG 413533

QY 918 GTGCACAATTATTCAGGAACACCCCAAAACCAAGTGAAGTGAATAGCATGAGAA 973
Db 413534 TAATGTAATAGCAACGCAATPAATAATCAATGATATAATAANGAGAAATAAGGA 413589

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RESULT 11
US-09-643-990A-1
; Sequence 1, Application US/09643990A
; Patent No. 6528289
; GENERAL INFORMATION:
; APPLICANT: Robert D. Fleischmann
; Mark D. Adams
; Owen White
; Hamilton O. Smith
; J. Craig Venter
; TITLE OF INVENTION: The Nucleotide sequence of
; the Haemophilus influenzae Rd Genome, Fragments
; Thereof, and Uses Thereof
; NUMBER OF SEQUENCES: 1
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville,
; STATE: MD
; COUNTRY: USA
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3 1/2 inch diskette
; COMPUTER: Dell Pentium
; OPERATING SYSTEM: MS DOS v6.22
; SOFTWARE: ASCII Text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/643,990A
; FILING DATE: 23-Aug-2000
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/487,429
; FILING DATE: 1995-06-07
; APPLICATION NUMBER: 08/426,787
; FILING DATE: 1995-04-21
; ATTORNEY/AGENT INFORMATION:
; NAME: Kenley K. Hoover
; REGISTRATION NUMBER: 40,302
; REFERENCE/DOCKET NUMBER: PB186P1C1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 301-610-5790

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; TELEFAX: 310-309-8439
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1830121 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-643-990A-1

Query Match          4.3%; Score 43; DB 4; Length 1830121;
Best Local Similarity 48.7%; Pred. No. 1.9;
Matches 115; Conservative 0; Mismatches 121; Indels 0; Gaps 0;

QY 738 AATTGTACAAAAAGTTTAAAGAAACTACATGTAAATCTTTGATAGTAAATAACTTGC 797
Db 413354 AAGTGTATACCAATATATAAGAGAGATGGAATAGTAAGATAGATAAATGCTGTT 413413

QY 798 CATTTCTTTATATGGAACGCAATTTTGGGTTGTTTAAAAATTTTAAACAGTTTATAAGAAA 857
Db 413414 TAAAGTTTAAATTTGATTTTCTATAGGTTGCTCAATTAATTAATAACTTTGTGATAGAAA 413473

QY 858 GAATTATAAGAAAAAGAAATACGCAATGGAAGTGTGAGTGAATAGCATGAGAA 917
Db 413474 AAATAATAGTTAATAATAACAACAATTAACATGTAATGCTTGAATCTGCTGCGCAG 413533

QY 918 GTGCACAATTATTCAGGAACACCCCAAAACCAAGTGAAGTGAATAGCATGAGAA 973
Db 413534 TAATGTAATAGCAACGCAATPAATAATCAATGATATAATAANGAGAAATAAGGA 413589

RESULT 12
US-08-232-463-14/c
; Sequence 14, Application US/08232463
; Patent No. 5670367
; GENERAL INFORMATION:
; APPLICANT: DORNER, F.
; APPLICANT: SCHEIFLINGER, F.
; APPLICANT: FALKNER, F. G.
; TITLE OF INVENTION: RECOMBINANT FOWLPOX VIRUS
; NUMBER OF SEQUENCES: 52
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Foley & Lardner
; STREET: 1800 Diagonal Road, Suite 500
; CITY: Alexandria
; STATE: VA
; COUNTRY: USA
; ZIP: 22313-0299
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/232,463
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/07/935,313
; FILING DATE:
; APPLICATION NUMBER: EP 91 114 300.6
; FILING DATE: 26-AUG-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: BENT, Stephen A.
; REGISTRATION NUMBER: 29,768
; REFERENCE/DOCKET NUMBER: 30472/114 IMMU
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703)836-9300
; TELEFAX: (703)683-4109
; TELEX: 899149
; INFORMATION FOR SEQ ID NO: 14:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7218 base pairs

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; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: linear
; IMMEDIATE SOURCE:
; CLONE: PTZpt-F1s
US-08-232-463-14

Query Match 4.3%; Score 42.8; DB 1; Length 7218;
Best Local Similarity 15.0%; Pred. No. 0.31;
Matches 41; Conservative 118; Mismatches 115; Indels 0; Gaps 0;

QY 701 TGTGTTTACAAATGGCAGGTTTACACGCTATGCAATGTACAAAAGATTATAGA 760
Db 1518 TATTGAGTTTCAAAAACGGCATGAGGCATCACTGTAATTAACCTATCTGCAAGTAGT 1459

QY 761 AAACCTACATGTAATCTTGTAGTAAATAAATCACTTGCATTTCTTTATATGGAACGCATT 820
Db 1458 TAAAGAGATAGAGAAATTTGGTACRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1399

QY 821 TTGGGTTGTTTAAATTTATAACAGTTATTAAGAAAGAAATTAAGGAAAAGAAAT 880
Db 1398 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1339

QY 881 AACGCAATGGCAAGTGGTGAAGCTGCAACTCAGCTGTGCACAAATTATCAGGAACACC 940
Db 1338 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1279

QY 941 CAAACCAAGTACGAGTGAAGATAGCATGAGAAG 974
Db 1278 RRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRRR 1245

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RESULT 13
US-08-487-826B-13
; Sequence 13, Application US/08487826B
; Patent No. 5993827
; GENERAL INFORMATION:
; APPLICANT: Sim, Kim L.
; APPLICANT: Chitnis, Chetan
; APPLICANT: Miller, Louis H.
; APPLICANT: Peterson, David S.
; APPLICANT: Su, Xin-zhaun
; APPLICANT: Wellens, Thomas E.
; TITLE OF INVENTION: BINDING DOMAINS FROM PLASMODIUM VIVAX
; NUMBER OF SEQUENCES: 45
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Knodbe Martens Olson & Bear
; STREET: 620 Newport Center Drive 16th Floor
; CITY: Newport Beach
; STATE: California
; COUNTRY: US
; ZIP: 92660
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/487,826B
; FILING DATE: 10-SEP-1993
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Israelson, Ned
; REGISTRATION NUMBER: 29,655
; REFERENCE/DOCKET NUMBER: NIH121.001CP1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 235-9550
; TELEFAX: (619) 235-0176
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 19124 base pairs
; TYPE: nucleic acid

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; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: CDNA
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
US-08-487-826B-13

Query Match 4.3%; Score 42.4; DB 2; Length 19124;
Best Local Similarity 50.8%; Pred. No. 0.54;
Matches 127; Conservative 0; Mismatches 121; Indels 2; Gaps 1;

QY 640 ATATTTTTCGAGGGTAGTGGTATTATATCTGTAGATTTCAGTAGTTCTTAAGTCGTGTTA 699
Db 4746 ATATATATATCATATGTAGTCATAGTGTCAATGAATATAAATATCGGTATATTATATTA 4805

QY 700 TTGTTTGTAAACAATGGCAGGTTTTCACGCTATGCAATTTGTACAAAAGATTATAG 759
Db 4806 TTGTATATATTAATAAGTAACACAGACATTTATATATATATATATATATATATATATAT 4865

QY 760 AAAACTACATGTAAATCTTGTATAGCTA--AATACTTGCCTATTTCTTTATATGGAACGC 817
Db 4866 ATATTTTGTATATATATATATATATATATATATATATATATATATATATATATAT 4925

QY 818 ATTTTGGGTTGTTTAAAAATTTATACAGTTTAAAGAAAGAAATTTAAAGGAAAAAGAA 877
Db 4926 AATTTTGTATATGATATAGTTATATAGTTAAAAAACAAGAAACAAATGGA 4985

QY 878 AATAACGCCAA 887
Db 4986 AAGCATAAAA 4995

```

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RESULT 14
US-09-091-432-3
; Sequence 3, Application US/09091432
; Patent No. 5981837
; GENERAL INFORMATION:
; APPLICANT: Chapelle, Clint
; TITLE OF INVENTION: A Method For Regulation Of Plant Lignin Composition
; FILE REFERENCE: 7024-325
; CURRENT APPLICATION NUMBER: US/09/091,432
; CURRENT FILING DATE: 1998-06-18
; EARLIER APPLICATION NUMBER: PCT/US96/20094
; EARLIER FILING DATE: 1996-12-19
; EARLIER APPLICATION NUMBER: US 60/009,119
; EARLIER FILING DATE: 1995-12-22
; EARLIER APPLICATION NUMBER: US 60/013,388
; EARLIER FILING DATE: 1996-03-14
; NUMBER OF SEQ ID NOS: 3
; SOFTWARE: Microsoft Word 2.0C
; SEQ ID NO 3
; LENGTH: 5156
; TYPE: DNA
; ORGANISM: Arabidopsis thaliana
US-09-091-432-3

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Query Match 4.2%; Score 42; DB 2; Length 5156;
Best Local Similarity 48.7%; Pred. No. 0.44;
Matches 114; Conservative 0; Mismatches 120; Indels 0; Gaps 0;

QY 559 AACCTCTGATTCCGCGCATGGTTTATATAATATTATATATATATATATATATATATAT 618
Db 1156 AAGTCTTGTGTAAAGCACTGTATCTAAATTTGTGATACGACAAATATATATATAT 1215

QY 619 GAGTTTACAAGAAATAAATCTATATTTTGTGAAGGGTAGTGGTATATATATCTAGTATT 678
Db 1216 GGACCTTCAAGTATTTATAGTATCTCTGTCTAAGATGCACAGGTATTCAGTAATAGTA 1275

QY 679 CAGTAGTTTCTAAGTCTGTTATTGTTTGTAAATGCGAGGTTTACACGCTCTATGCA 738
Db 1276 AAATAATCTTGTATATATATATATCTAATAGTAACCTTGTCTTAACCTAAATGAG 1335

QY 739 ATTGTACAAAAAGTTTATAGAAAACTACATGTAATAAATCTTTGATAGCTAAATAA 792

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Search completed: April 6, 2004, 14:14:25
Job time : 118.764 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 03:49:50 ; Search time 3674.56 Seconds
(without alignments)
8094.223 Million cell updates/sec

Title: US-09-966-264D-2
Perfect score: 996
Sequence: 1 gtggttgattgattagtaaa.....gtgttgattgatttaatt 996

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 27513289 seqs, 14931090276 residues

Total number of hits satisfying chosen parameters: 55026578

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

EST: *
1: em_estba:*
2: em_esthum:*
3: em_estin:*
4: em_estmu:*
5: em_estov:*
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7: em_estro:*
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17: em_gss_hum:*
18: em_gss_inv:*
19: em_gss_pln:*
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23: em_gss_mus:*
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25: em_gss_rod:*
26: em_gss_phg:*
27: em_gss_vrl:*
28: gb_gss1:*
29: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	459.2	46.1	525	28	AQ679243 HS 5417 A
2	443.2	44.5	472	28	AQ583263 RBCI-11-4
3	410.8	84.4	41.2	844	11 BG706268
4	375.8	37.7	4437	11	AK036936 Mus muscu

5	370.4	37.2	439	14	CB750007
6	354	35.5	924	13	BQ27942
7	342.8	34.4	629	9	AL712236
8	342.8	34.4	629	13	BX509925
9	296	29.7	633	13	BX337294
10	291.2	29.2	531	12	BI281078
11	286.2	28.7	854	13	BX226117
12	282.8	28.4	739	13	BX368789
13	282.6	28.4	500	9	AI008807
14	281.2	28.2	576	12	BG795230
15	278.6	28.0	421	13	BY635064
16	273.8	27.5	658	13	BY352875
17	271.6	27.3	308	14	N75050
18	270.8	27.2	338	13	BX354233
19	269	27.0	440	10	BB819684
20	266.2	26.7	687	14	W05777
21	264.2	26.5	929	13	BU274687
22	263.2	26.4	778	13	BU309070
23	260.2	26.1	492	14	CB725360
24	245.4	24.6	999	13	EX419180
25	245	24.6	648	13	BU402013
26	244.8	24.6	739	9	AUI38648
27	237.2	23.8	826	13	BU104673
28	233.8	23.5	1201	13	BX463891
29	231.2	23.2	859	13	BU300671
30	229.2	23.0	928	13	BU107395
31	221.2	22.2	927	13	BU139194
32	217.8	21.9	888	14	CF553038
33	215	21.6	277	10	AW861957
34	213.8	21.5	275	10	AW604756
35	213.8	21.5	284	10	AW858457
36	208	20.9	234	14	N86611
37	202.8	20.4	619	10	AW71158
38	202.2	20.3	271	10	AW858578
39	195.6	19.6	886	13	EX760461
40	195.4	19.6	851	13	BX725372
41	194.8	19.6	628	13	BX693509
42	193.8	19.5	677	13	BX855319
43	192.8	19.4	618	9	AL793265
44	191	19.2	537	12	BU090827
45	186.6	18.7	351	13	BY673194

ALIGNMENTS

RESULT 1	AQ679243	525 bp	DNA	linear	GSS 25-JUN-1999
LOCUS	HS5417	Al B06 T7A	RBCI-11	Human Male	BAC Library Homo sapiens
DEFINITION	genomic clone Plate=993	Col=11	Row=C	genomic survey	sequence.
ACCESSION	AQ679243				
VERSION	AQ679243.1	GI:5228047			
KEYWORDS	GSS.				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
REFERENCE	1 (bases 1 to 525)				
AUTHORS	Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and Hood,L.				
TITLE	Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome				
JOURNAL	Proc. Natl. Acad. Sci. U.S.A.	96 (17)			9739-9744 (1999)
MEDLINE	9930589				
PUBMED	10449764				
COMMENT	Contact: Mahairas GG, Wallace JC, Hood L High Throughput Sequencing Center University of Washington 401 Queen Anne Avenue North, Seattle, WA 98109, USA Tel: (206) 616-3618 Fax: (206) 616-3887				

CB750007 AMGNNUC:S
BQ27942 AGENCOURT
AL712236 DKFZp686J
BX509925 DKFZp686J
BU337294 60351575J
BI281078 UI-R-DD0-
BU226117 603947264
BU368789 603596716
AI008807 EST03258
BG795230 UTSW SM30
BY635064 BY635064
BU352875 603525781
N75050 248907.81
BU352423 603530318
BB819684 BB819684
W05777 248907.r1
BU274687 603531284
BU309070 603538118
CB725360 AMGNNUC:N
EX419180 EX419180
BU402013 604138944
AUI38648 AUI38648
BU104673 603004172
BX463891 BX463891
BU300671 603734536
BU107395 603110749
BU139194 603135045
CF553038 AGENCOURT
AW861957 CM3-CT034
AW604756 CM0-CT034
AW858457 CM3-CT034
N86611 J9202F Huma
AW71158 hn53c06.x
AW858578 CM3-CT034
EX760461 EX760461
BX725372 BX725372
BX693509 BX693509
BX855319 BX855319
AL793265 AL793265
BU090827 BU090827
BY673194 BY673194

Email: jwallace@u.washington.edu
 Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm) or from Research Genetics (info@resgen.com). BAC end Web Server: http://www.hnsc.washington.edu
 Plate: 993 row: C column: 11
 Seq primer: T7
 Class: BAC ends
 High quality sequence stop: 525.

FEATURES
 source
 1. 525
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /clone="Plate:993 Col=11 Row=C"
 /sex="male"
 /clone_lib="RPCI-11 Human Male BAC Library"
 /note="Vector: pBACe3.6; Site:1: EcoRI; Site:2: EcoRI; Male blood DNA was isolated from one randomly chosen donor and partially digested with a combination of EcoRI and EcoRI Methylase. Size selected DNA was cloned into the pBACe3.6 vector at EcoRI sites"

ORIGIN
 Query Match 46.1%; Score 459.2; DB 28; Length 525;
 Best Local Similarity 95.4%; Pred. No. 3.4e-72; Indels 1; Gaps 1;
 Matches 481; Conservative 0; Mismatches 22;
 QY 137 TCTATGGAATCTTTTGATATATATTTACATGGGAACCTGAATGTAGCTTGACATTTT 196
 Db 23 TCTCTCGGACTGCCNTGATATATATNTACATGGGAACCTGAATGTAGCTTGACATTTT 82
 QY 197 CCATGTAAACACCACTAGCTGATCCACATTAAGCTATCTATACACACACCTGTAAT 256
 Db 83 CCATGTAAACACCACTAGCTGATCCACATTAAGCTATCTATACACACACCTGTAAT 142
 QY 257 GGCCTTCATTATAGAGCTTTGCTTCTCTGGAACCTGGTGAAGAAATCAAACTTTGTTG 316
 Db 143 GGCCTTCATTATAGAGCTTTGCTTCTCTGGAACCTGGTGAAGAAATCAAACTTTGTTG 202
 QY 317 GTACACCTCGATGAGCTTCTGTGTTCTTCACCCAGAAATGGGAATGATTTCCAA 376
 Db 203 GTACACCTCGATGAGCTTCTGTGTTCTTCACCCAGAAATGGGAATGATTTCCAA 262
 QY 377 ATGGCAAGAACACAGATGATCTATCTATCTGACCTTTTGTAAAGTCTGCTTTCTTT 436
 Db 263 ATGGCAAGAACACAGATGATCTATCTATCTGACCTTTTGTAAAGTCTGCTTTCTTT 321
 QY 437 CTCCTTTGTTTCCAGGACACAAATGAGGAAGTCTTTTCCACATGGCAGATGATTGGCA 496
 Db 322 CTCCTTTGTTTCCAGGACACAAATGAGGAAGTCTTTTCCACATGGCAGATGATTGGCA 381
 QY 497 GAGCGATGAGTCCCTTAGTATCAGTCATGACAGATGAAGAGGACAGAAATGATTTT 556
 Db 382 GAGCGATGAGTCCCTTAGTATCAGTCATGACAGATGAAGAGGACAGAAATGATTTT 441
 QY 557 ACAATCTCTGATCCCGCATGTTTATATATTTATATATATATATATATATATATATAT 616
 Db 442 ACAATCTCTGATCCCGCATGTTTATATATATATATATATATATATATATATATATAT 501
 QY 617 AAGAGTTTACAGAAATTAATCTA 640
 Db 502 TAGAGTTTACAGAAATTAATCTA 525

RESULT 2
 A0583263
 LOCUS
 DEFINITION
 RPCI-11-451G14, TV RPCI-11 Homo sapiens genomic clone
 A0583263
 ACCESSION

VERSION
 A0583263.1 GI:5010373
 GSS.
 KEYWORDS
 SOURCE
 ORGANISM
 Homo sapiens (human)
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 472)
 Zhao,S., Adams,M.D., Nierman,W., Malek,J., de Jong,P. and
 Venter,J.C.
 Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
 Map Building
 Unpublished (1997)
 Contact: Shaying Zhao, William Nierman, Mark Adams
 Department of Eukaryotic Genomics
 the Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: hbeetig@ig.org
 Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genet cs (info@resgen.com). BAC end search page: http://www.tigr.org/tdb/hungen/bac_end_search/bac_end_search.html.
 Seq primer: T7
 Class: BAC ends.
 Location/Qualifiers
 1..472
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="GDB:7672957"
 /db_xref="taxon:9606"
 /clone="RPCI-11-451G14"
 /sex="Male"
 /cell_type="Lymphocytes"
 /clone_lib="RPCI-11"
 /note="Vector: pBACe3.6; Site:1: EcoRI; Site:2: EcoRI; RPCI11 Human Male BAC Library"

FEATURES
 source
 1..472
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="GDB:7672957"
 /db_xref="taxon:9606"
 /clone="RPCI-11-451G14"
 /sex="Male"
 /cell_type="Lymphocytes"
 /clone_lib="RPCI-11"
 /note="Vector: pBACe3.6; Site:1: EcoRI; Site:2: EcoRI; RPCI11 Human Male BAC Library"

ORIGIN
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 Best Local Similarity 96.2%; Pred. No. 2.5e-69; Indels 0; Gaps 0;
 Matches 454; Conservative 0; Mismatches 18;
 QY 172 AACCTGAATGTAGCTGACATTTTCCATGTAACACCACTAGCTGATCCCAACATTAG 231
 Db 1 AACCTGAATGTAGCTGACATTTTCCATGTAACACCACTAGCTGATCCCAACATTAG 60
 QY 232 CTGATCTAACAAACAAACCGTGAATGGCTTCAATTAAGGCTTTGCTTCTTCTGGAAA 291
 Db 61 CTGATCTAACAAACAAACCGTGAATGGCTTCAATTAAGGCTTTGCTTCTTCTGGAAA 120
 QY 292 CTGGTGAAGAAATCAAAACCTTTGTTGTACACCTCGATGACAGCTTCTGTGTTCTTCC 351
 Db 121 CTGGTGAAGAAATCAAAACCTTTGTTGTACACCTCGATGACAGCTTCTGTGTTCTTCC 180
 QY 352 CCGAATATGGGAATGATTTCCCAATGCAAGAAACAGAGTGATGCTATCTATCTGCA 411
 Db 181 CCGAATATGGGAATGATTTCCCAATGCAAGAAACAGAGTGATGCTATCTATCTGCA 240
 QY 412 CCTTTTGTAAAGTCTGCTTTCTTTCTTTTCCAGGACACAATGAGGAAGTCTT 471
 Db 241 CCTTTTGTAAAGTCTGCTTTCTTTCTTTTCCAGGACACAATGAGGAAGTCTT 300
 QY 472 TTCACATGGCAGATGATTTGGCAGAGGATGGAGTCTCTAGTATCAGTCATGACAGAT 531
 Db 301 TTCACATGGCAGATGATTTGGCAGAGGATGGAGTCTCTAGTATCAGTCATGACAGAT 360
 QY 532 GAAGAGGACAGAAATATTTTCAACTCTGATTCCTGATTCCTGATTCCTGATTCCTGATTC 591
 Db 361 GAAGAGGACAGAAATATTTTCAACTCTGATTCCTGATTCCTGATTCCTGATTCCTGATTC 420

Hori, F., Imotani, K., Iehii, Y., Itoh, M., Kagawa, I., Kasukawa, T., Kato, H., Kawai, J., Kojima, Y., Kondo, S., Kono, H., Kouda, M., Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, M., Nakamura, M., Nishi, K., Nomura, K., Numazaki, R., Ono, M., Ohsato, N., Okazaki, Y., Saito, R., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N., Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Tagami, M., Tagawa, A., Takahashi, F., Takaku-Akahira, S., Takeda, Y., Tanaka, T., Tomaru, A., Toya, T., Yasunishi, A., Muramatsu, M., and Hayashizaki, Y.

Direct Submission
Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of Physical and Chemical Research (RIKEN), Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), RIKEN Yokohama Institute, 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan [E-mail: genome-res@gsc.riken.go.jp, URL: http://genome.gsc.riken.go.jp/, Tel: 81-45-503-9222, Fax: 81-45-503-9216]

CDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Please visit our web site for further details.
URL: http://genome.gsc.riken.go.jp/
URL: http://fantom.gsc.riken.go.jp/
Location/Qualifiers
1. .4437
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="PANTOM_DB:9930028B14"
/db_xref="MGI:2401399"
/db_xref="taxon:10090"
/clone="9930028B14"
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/tissue_type="vagina"
/clone_lib="RIKEN full-length enriched mouse cDNA library"
/dev_stage="adult"
29. .1988
/note="dystrophin, muscular dystrophy (MGD|MGI:94909, GB|NM_007868, evidence: BLASTN, 100%, match=99); putative"
polyA_signal
4417. .4422
/note="putative"
polyA_site
4437
/note="putative"

misc_feature
29. .1988
/note="dystrophin, muscular dystrophy (MGD|MGI:94909, GB|NM_007868, evidence: BLASTN, 100%, match=99); putative"

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/note="putative"

polyA_site
4437
/note="putative"

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Query Match 37.7%; Score 375.8; DB 11; Length 4437;
Best Local Similarity 96.2%; Pred. No. 1e-57;
Matches 407; Conservative 0; Mismatches 12; Indels 4; Gaps 2;
QY 450 AGGACACATGATGAGGAAGTCCTTTCCACATGCGCAGATGTTGGCAGAGCGATGGAGTC 509
DB 1869 AGGACACATGATGAGGAAGTCCTTTCCACATGCGCAGATGTTGGCAGAGCGATGGAGTC 1928
QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGCAATAATGTTTACAACTCCTGATT 569
DB 1929 CTTAGTTCAGTCATGACAGATGAAGAAGGAGCAGCAATAATGTTTACAACTCCTGATT 1988
QY 570 CCGCATGTTTATATATATTCATACACAAAGAGGATTAGACAGTAAGAGTTTACAG 629
DB 1989 CCGCATGTTTATATATATTCATACACAAAGAGGATTAGACAGTAAGAGTTTACAG 2048
QY 630 AAAT-AAAATCTATATTTTGTGAAGGAGTGTGTTATATCTGATGATTTCAGTAGTTTC 688
DB 2049 AAATAAATCTATATTTTGTGAAGGAGTGTGTTATCTGATGATTTCAGTAGTTTC 2108
QY 689 TAAGTCTGTTATGTTTGTGTTAAACAAAGCGAGTTTACAGTCTATGCAATTTGACAAA 748
DB 2109 TAAGTCTGTTATGTTTGTGTTAAACAAAGCGAGTTTACAGTCTATGCAATTTGACAAA 2168
QY 749 AAAGTATTAAGAAACTACATGTAATACTTGTATAGCTAAATAAATCTGCCATTTCTTTAT 808

DB 2169 AAAGTATTAAGAAA---ACAATGTAATACTTGTAGTAAATAAATCTGCCATTTCTTTAT 2225
QY 809 ATGGACCGCATTTGGGTGTTTAAAAATTTATAACAGTTTATAAAGAAGAAATTAAGAAG 868
DB 2226 ATGGACCGCATTTGGGTGTTTAAAAATTTATAACAGTTTATAAAGAAGAAATTTGAAGT 2285
QY 869 GAA 871
DB 2286 AAA 2288

RESULT 5
CB750007
LOCUS
DEFINITION
AMGNNUC:SRPB2-00166-B9-A.srp2 (10220) Rattus norvegicus cDNA clone
srp2-00166-b9 5', mRNA sequence.
CB750007 439 bp mRNA linear EST 11-APR-2003
CB750007.1 GI:29817309
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Rattus norvegicus (Norway rat)
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 439)
Angen EST Program.
Angen Rat EST Program
Unpublished (2003)
Contact: Dan Fitzpatrick
Angen, Inc
One Angen Center Drive, Thousand Oaks, CA 91320-1799, USA
Tel: 805 447-4881
Plate: 00166 row: b column: 9.
Location/Qualifiers
1. .439
/organism="Rattus norvegicus"
/mol_type="mRNA"
/db_xref="taxon:10116"
/clone="srp2-00166-b9"
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/clone_lib="srp2 (10220)"
/note="Vector: pSPORT1; Site 1: SalI; Site 2: NotI; rat
prostate normalized double selected poly(A+) mRNA size
fraction > 1 kb"

FEATURES
source
1. .439
/organism="Rattus norvegicus"
/mol_type="mRNA"
/db_xref="taxon:10116"
/clone="srp2-00166-b9"
/tissue_type="prostate tissue"
/clone_lib="srp2 (10220)"
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prostate normalized double selected poly(A+) mRNA size
fraction > 1 kb"

ORIGIN
Query Match 37.2%; Score 370.4; DB 14; Length 439;
Best Local Similarity 95.4%; Pred. No. 2.4e-56;
Matches 393; Conservative 0; Mismatches 16; Indels 3; Gaps 1;
QY 460 GTAGGAAGTCCTTTCCACATGCGCAGATGATTGGGCGAGAGCGATGGAGTCCTTAGTATCA 519
DB 12 GTAGGAAGTCCTTTCCACATGCGCAGATGATTGGGCGAGAGCGATGGAGTCCTTAGTATCA 71
QY 520 GTCATGACAGATGAAGAAGGAGCAGCAATAATGTTTACAACTCCTGATCCCGCATGTT 579
DB 72 GTCATGATAGATGAAGAAGGAGCAGCAATAATGTTTACAACTCCTGATCCCGCATGTT 131
QY 580 TTTTATAATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAAGAAATAAATCT 639
DB 132 TTTTATAATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAAGAAATAAATCT 191
QY 640 ATATTTTGTGAAGGAGTGTGTTATATCTAGTATTTAGTATTTTCAAGTCTGTTA 699
DB 192 ATATTTTGTGAAGGAGTGTGTTATCTAGTATTTTCAAGTCTGTTA 251
QY 700 TTTTGTGTTTAAAGTGGCAGGTTTACACGCTCTGCAATTTGTCACAAAAGTTTAAAG 759
DB 252 TTTGTTTGTAAAGTGGCAGGTTTACACGCTCTGCAATTTGTCACAAAAGTTTAAAG 311
QY 760 AAAACTACATGTAATAATCTTGTAGTAAATAAATCTGCCATTTCTTTATATGGAACGAT 819

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Db      312 AAA--ACATGTAATCTTGATAGCTAAATAACTTCCATTTCTTTATATGGAACGCAT 368
Qy      920 TTGGGTTGTTTAAATTTTATACAGTTTATAAGAAAGAAATTAAGGAA 871
Db      369 TTGGGTTGTTTAAATTTTATACAGTTTATAAGAAAGAAATTAAGGAA 420

RESULT 6
BQ927942      BQ927942      924 bp      mRNA      linear      EST 20-AUG-2002
LOCUS      AGENCOURT_8858741 NCI_CGAP Mam2 Mus musculus cDNA clone
DEFINITION      IMAGE:6441872 5', mRNA sequence.
ACCESSION      BQ927942
VERSION      BQ927942.1 GI:22342973
KEYWORDS      EST.
SOURCE      Mus musculus (house mouse)
ORGANISM      Mus musculus
REFERENCE      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS      Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
TITLE      NIH-MGC http://mgi.nci.nih.gov/.
JOURNAL      National Institutes of Health, Mammalian Gene Collection (MGC)
COMMENT      Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Gilbert Smith, Ph.D.,
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM13965 row: m column: 09
High quality sequence start: 10
High quality sequence stop: 552.

FEATURES             Location/Qualifiers
     source            1..924
     organism="Mus musculus"
     mol_type="mRNA"
     strain="FVB/N-3"
     db_xref="taxon:10090"
     clone="IMAGE:6441872"
     tissue_type="tumor, biopsy sample"
     dev_stage="5 months"
     lab_host="DH10B"
     clone_lib="NCI CGAP Mam2"
     note="Organ: mammary; Vector: pCMV-SPOK16; Site_1: SalI;
     Site_2: NotI; Cloned unidirectionally. Primer: Oligo dt.
     Library constructed by Life Technologies. Investigator
     providing samples: Gilbert Smith, NIH"

ORIGIN
Query Match      35.5%; Score 354; DB 13; Length 924;
Best Local Similarity 95.3%; Pred. No. 1.5e-53;
Matches 387; Conservative 0; Mismatches 15; Indels 4; Gaps 2;

Qy      467 GTCTTTCCACATGCCAGATGTTGGCAGAGCGATGGAGTCTCTAGTATCAGTCATGA 526
Db      33  GGCCTTTTCCCATGCCAGATGTTGGCAGAGCGATGGAGTCTCTAGTATCAGTCATGA 92
Qy      527 CAGATGAAGAGGAGCAGAAATAATGTTTCAACTCTGATTCGCCGATGGTTTTATA 586
Db      93  CAGATGAAGAGGAGCAGAAATAATGTTTCAACTCTGATTCGCCGATGGTTTTATA 152
Qy      587 ATATTTCATACAAAGAGGATTACACAGTAGAGTTTACAGAAAT-AAATCTATATTT 645
Db      153 ATATTTCGTAACAAGAGGATTACACAGTAGAGTTTACAGAAATATAATCTATATTT 212
Qy      646 TTGTGAAGGGTAGTGGTATTATPACTGTAGATTTCAGTAGTTTCTAAGTCTGTTATGTTT 705
Db      213 TTGTGAAGGGTAGTGGTACTATPACTGTAGATTTCAGTAGTTTCTAAGTCTGTTATGTTT 272
Qy      706 TGTAAACATGCGCAGTTTACACGCTCTATGCAATGTACAAAAGTTTAAGAAACT 765

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Db      273 TGTAAACATGCGCAGTTTACACGCTCTATGCAATTTGTACAAAAGTTTAAAGAAA--- 329
Qy      766 ACATGTAATCTTGATAGCTAAATAACTTCCATTTCTTTATATGGAACGCATTTGGG 825
Db      330 ACATGTAATCTTGATAGCTAAATAACTTCCATTTCTTTATATGGAACGCATTTGGG 389
Qy      826 TTGTTTAAAAATTTATAACAGTTTATAAGAAAGAAATTAAGGAA 871
Db      390 TTGTTTAAAAATTTATAACAGTTTATAAGAAAGAAATTTGTAATACTAA 435

RESULT 7
AL712236      629 bp      mRNA      linear      EST 04-SEP-2003
LOCUS      DKFZp686J0287 r1.686 (synonym: hlcc3) Homo sapiens cDNA clone
DEFINITION      DKFZp686J0287 5', mRNA sequence.
ACCESSION      AL712236
VERSION      AL712236.2 GI:32026719
KEYWORDS      EST.
SOURCE      Homo sapiens (human)
ORGANISM      Homo sapiens
REFERENCE      Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS      Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
TITLE      1 (bases 1 to 629)
JOURNAL      Poustka, A., Albert, R., Moosmayer, P., Schupp, I., Wellenreuther, R.,
COMMENT      Wewes, H.W., Weil, B., Amid, C., Osanger, A., Pobo, G., Han, M. and
                Wiemann, S.
                EST (Poustka, A., Albert, R., Moosmayer, P., Schupp, I.,
                Wellenreuther, R., et al.)
                Unpublished (2003)
                On Mar 22, 2002 this sequence version replaced gi:19695591.
                Contact: MIPS
                MIPS
                Ingolstaedter Landstr.1, D-85764 Neuherberg, Germany
                This is the 5' sequence of the clone insert
                Clone from S. Wiemann, Molecular Genome Analysis, German Cancer
                Research Center (DKFZ); Email s.wiemann@dkfz-heidelberg.de;
                sequenced by DKFZ (German Cancer Research Center,
                Heidelberg/Germany) within the cDNA sequencing consortium of the
                German Genome Project.
                No sl sequence available.
                This clone (DKFZp686J0287) is available at the RZPD in Berlin.
                Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059
                Berlin-Charlottenburg, GERMANY; Email: clone@rzpd.de.

FEATURES             Location/Qualifiers
     source            1..629
     organism="Homo sapiens"
     mol_type="mRNA"
     db_xref="taxon:9606"
     clone="DKFZp686J0287"
     dev_stage="adult"
     lab_host="DH10B"
     clone_lib="686 (synonym: hlcc3)"
     note="Vector: pTriplex2; Site_1: SfiI; Site_2: SfiI;
     cDNA-collection"

ORIGIN
Query Match      34.4%; Score 342.8; DB 9; Length 629;
Best Local Similarity 98.0%; Pred. No. 1.7e-51;
Matches 347; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

Qy      518 CAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTACAACTCCTGATTCGCCCATG 577
Db      1  CAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTACAACTCCTGATTCGCCCATG 60
Qy      578 GTTTTTATATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAGAAATAAAT 637
Db      61  GTTTTTATATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAGAAATAAAT 120
Qy      638 CTATATTTTGTGAAGGGTAGTGGTATTACTGTAGATTTCAGTAGTTTCTAAGTCTCT 697
Db      121 CTATATTTTGTGAAGGGTAGTGGTATTACTGTAGATTTCAGTAGTTTCTAAGTCTCT 180

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QY 698 TATTGTTTGTAAATGGAGGTTTACAGTCTATGCAATTTGTACAAAAGTTTATA 757
DB 181 TATTGTTTGTAAATGGAGGTTTACAGTCTATGCAATTTGTACAAAAGTTTATA 240
QY 758 AGAAACTACATGTAATCTTGTAGTAAATTAACCTTGCCTTTCTTTATATGGAACGC 817
DB 241 AGAAACTACATGTAATCTTGTAGTAAATTAACCTTGCCTTTCTTTATATGGAACGC 300
QY 818 ATTTGGTGTGTTTAAAAATTTTAAACAGTTTATAAGAAAGTAATATAAGGAA 871
DB 301 ATTTGGTGTGTTTAAAAATTTTAAACAGTTTATAAGAAAGTAATATAAGGAA 354

RESULT 8
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LOCUS DKFZP686J2487_r1 686 (synonym: hicc3) Homo sapiens cDNA clone
DEFINITION DKFZP686J2487 5', mRNA sequence.
ACCESSION BX509925
VERSION BX509925
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 629)
AUTHORS Poustka,A., Albert,R., Moosmayer,P., Schupp,I., Wellenreuther,R.,
Mewes,H.W., Weil,B., Amid,C., Osanger,A., Fobo,G., Han,M. and
Wiemann,S.
TITLE EST (Poustka,A., Albert,R., Moosmayer,P., Schupp,I.,
Wellenreuther,R., et al.)
JOURNAL Unpublished (2003)
COMMENT Contact: MIPS
MIPS Ingolstaedter Landstr.1, D-85764 Neuherberg, Germany
This is the 5' sequence of the clone insert
Clone from S. Wilmann, Molecular Genome Analysis, German Cancer
Research Center (DKFZ); Email s.wilmann@dkfz-heidelberg.de;
Heidelberg/Germany) within the cDNA sequencing consortium of the
German Genome Project.
No s1 sequence available.
This clone (DKFZP686J2487) is available at the RZPD in Berlin.
Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059
Berlin-Charlottenburg, GERMANY; Email: clone@rzpd.de.
Location/Qualifiers
1. .629
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
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/dev_stage="adult"
/lab_host="DH10B"
/note="Vector: pTriplex2; Site_1: sfliA; Site_2: sfliB;
cDNA-collection"

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Query Match 34.4%; Score 342.8; DB 13; Length 629;
Best Local Similarity 98.0%; Pred. No. 1.7e-51;
Matches 347; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 518 CAGTCATGACAGATGAAGAGGAGCAGATAATGTTTACAACTCCTGATTCGCGCATG 577
DB 1 CAGTCATGACAGATGAAGAGGAGCAGATAATGTTTACAACTCCTGATTCGCGCATG 60
QY 578 GTTTTATATATATCTATACACAAAGAGGATTAGACAGTAAGAGTTTACAGAAATAAT 637
DB 61 GTTTTATATATATCTATACACAAAGAGGATTAGACAGTAAGAGTTTACAGAAATAAT 120
QY 638 CTATATTTTGTCAAGGTTAGTGTATATATCTGTAGATTTCAGTAGTTTCTAAGTCTGT 697

DB 121 CTATATTTTGTGAAGGTTAGTGGTATTATATCTGTAGATTTCAGTAGTTTCTAAGTCTGT 180
QY 698 TATTGTTTGTAAATGGAGGTTTACAGTCTATGCAATTTGTACAAAAGTTTATA 757
DB 181 TATTGTTTGTAAATGGAGGTTTACAGTCTATGCAATTTGTACAAAAGTTTATA 240
QY 758 AGAAACTACATGTAATCTTGTAGTAAATTAACCTTGCCTTTCTTTATATGGAACGC 817
DB 241 AGAAACTACATGTAATCTTGTAGTAAATTAACCTTGCCTTTCTTTATATGGAACGC 300
QY 818 ATTTGGTGTGTTTAAAAATTTTAAACAGTTTATAAGAAAGTAATATAAGGAA 871
DB 301 ATTTGGTGTGTTTAAAAATTTTAAACAGTTTATAAGAAAGTAATATAAGGAA 354

RESULT 9
BU337294 633 bp mRNA linear EST 28-NOV-2002
LOCUS 60351575F1 CSEQCHN66 Gallus gallus cDNA clone Chest455a15 5', mRNA
DEFINITION sequence.
ACCESSION BU337294
VERSION BU337294
KEYWORDS EST.
SOURCE Gallus gallus (chicken)
ORGANISM Gallus gallus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
Phasianinae; Gallus.
REFERENCE 1 (bases 1 to 633)
AUTHORS Boardman,P.E., Sanz-Ezquerro,J., Overton,I.M., Burt,D.W., Bosch,E.,
Fong,W.T., Tickle,C., Brown,W.R.A., Wilson,S.A. and Hubbard,S.J.
TITLE A Comprehensive Collection of Chicken cDNAs
JOURNAL Curr. Biol. 12 (22), 1965-1969 (2002)
MEDLINE 22335534
PUBMED 12445392
COMMENT Contact: Simon Hubbard
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University of Manchester Institute of Science and Technology
(UMIST)
PO Box 88, Manchester, M60 1QD, UK
Tel: 01612008930
Fax: 01612360409
Email: Simon.Hubbard@umist.ac.uk.
Location/Qualifiers
1. .633
/organism="Gallus gallus"
/mol_type="mRNA"
/strain="White Leghorn, Hisex"
/db_xref="taxon:9031"
/clone="Chest455a15"
/dev_stage="16 day embryo"
/lab_host="DH10B"
/clone_lib="CSEQCHN66"
/note="Organ: brain; Vector: pBluescript II KS(+); Site_1:
EcoRI; Site_2: NotI; this normalized library was
constructed from 1 million independent clones. cDNA
synthesis was initiated using an oligo(dT) primer, using
methylated C in the first strand synthesis reaction.
Following this first strand reaction, double-stranded cDNA
was blunted, ligated to NotI adapters, digested with
EcoRI. Size-selected, and cloned into the NotI and EcoRI
compatible sites of a custom modified MCS of the
pBluescript (KS+) vector. The library was normalized in 2
rounds using conditions adapted from Soares et al., PNAS
(1994) 91: 9228-9232 and Bonaldo et al., Genome Research 6
(1996): 791, except that a significantly longer
reannealing hybridization was used."

ORIGIN
Query Match 29.7%; Score 296; DB 13; Length 633;
Best Local Similarity 87.2%; Pred. No. 3.8e-43;
Matches 396; Conservative 0; Mismatches 45; Indels 13; Gaps 6;

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Fax: 01612360409
Email: Simon.Hubbard@umist.ac.uk.
Location/Qualifiers

FEATURES

source
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/mol_type="mRNA"
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constructed from 1 million independent clones. cDNA
synthesis was initiated using an oligo(dT) primer, using
methylated C in the first strand synthesis reaction.
Following this first strand reaction, double-stranded cDNA
was blunted, ligated to NotI adapters, digested with
EcoRI, size-selected, and cloned into the NotI and EcoRI
compatible sites of a custom modified MCS of the
pBluescript (KS+) vector. The library was normalized in 2
rounds using conditions adapted from Soares et al., PNAS
(1994) 91: 9228-9232 and Bonaldo et al., Genome Research 6
(1996): 791, except that a significantly longer
reannealing hybridization was used."

ORIGIN

Query Match 28.7%; Score 286.2; DB 13; Length 854;
Best Local Similarity 86.3%; Pred. No. 1.8e-41;
Matches 389; Conservative 0; Mismatches 48; Indels 14; Gaps 6;
QY 437 CTCCTTGTTCAGGACACATGATGAGNAGCTTTTCCACATGCGCAGATGTTGGCA 496
DB 5 CTCGAGGTTCCAGAGGCCCAATGATGAGGAGCTTTTCCACATGCGCAGATGTTGGAA 64
QY 497 GAGCGATGAGTCCCTAGTATCAGTCATGACAGATGAAGAGGAGCAGAAT-----AAA 550
DB 65 GAGCGATGGAACCTTAGTGACGGTGATGACGACGACAAAGGTGTAGATAGCAAGATG 124
QY 551 TGTTTTAAAC--TCCTGATCCCGCATGTTTATATATATATATATATATATATATATATAT 609
DB 125 TGTTTTAAACCTCAGCGCTCCCGCATGTTTATATATATATATATATATATATATATATAT 184
QY 610 AGACATGAAGATTACAG--AAATAAATCTATATTTTGTCAAGGCTAGTGGTATTATA 668
DB 185 AGACTGTAAGATTACAAAGAAAACAAATCTACATTTTGTGAAGGCTAGTGGTACTATA 244
QY 669 CTGTAGATTTCAGTAGTTT--CTAAGTCTGTTATTTGTTTGTAAACAATGCGAGTTTAC 727
DB 245 CTGTAGATTTCAGTAGTTTCTCCTAAGTCTGTTTCTGTTTGTAAACAATGCGAGTTTAC 304
QY 728 AGCTCTACGATTTACAAAAGTTATAGAACAATACATGTAATCTTTGATAGCTA 787
DB 305 AGCTCTACGATTTACAAA--TTATAAGAAACATCAATGTAATCTTTGATAGCTG 362
QY 788 AATAACTTGCATTTCTTTATATGGAACGCAATTTGGGTTGTTT---AAAAATTTATAAC 844
DB 363 AATAACTTGCATTTCTTTATATGGAACGCAATTTGGGTTGTTTAAAAAATTTATAAC 422
QY 845 AGTTATAAGAAAGATTTAAGGAAAAAG 875
DB 423 AGTTATAAGAACAGATTGTAACCTAAAAAG 453

RESULT 12

BU368789 739 bp mRNA linear EST 28-NOV-2002
LOCUS 603596716F1 CSEQCHN73 Gallus gallus cDNA clone ChEST56503 5', mRNA
DEFINITION sequence.
ACCESSION BU368789

VERSION KEYWORDS SOURCE ORGANISM

BU368789.1 GI:25876790
EST.
Gallus gallus (chicken)
Gallus gallus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
Phasianinae; Gallus.
1 (bases 1 to 739)
Boardman, P.E., Sanz-Ezquerro, J., Overton, I.M., Burt, D.W., Bosch, E.,
Fong, W.T., Tickle, C., Brown, W.R.A., Wilson, S.A. and Hubbard, S.J.
A Comprehensive Collection of Chicken cDNAs
Curr. Biol. 12 (22), 1965-1969 (2002)
22335534
MEDLINE
12445392
PUBMED

REFERENCE AUTHORS

TITLE

JOURNAL

MEDLINE

PUBMED

COMMENT

Contact: Simon Hubbard
Department of Biochemical Sciences
University of Manchester Institute of Science and Technology
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Tel: 01612008930
Fax: 01612360409
Email: Simon.Hubbard@umist.ac.uk.
Location/Qualifiers

FEATURES source

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/organism="Gallus gallus"
/mol_type="mRNA"
/strain="Compton Line 151"
/db_xref="taxon:9031"
/clone="ChEST56503"
/sex="Female"
/tissue_type="not cerebrum or cerebellum"
/dev_stage="adult"
/lab_host="DH10B"
/clone_lib="CSEQCHN73"
/note="Organ: brain; Vector: pBluescript II KS(+); Site_1:
EcoRI; Site 2: NotI; This normalized library was
constructed from 1 million independent clones. cDNA
synthesis was initiated using an oligo(dT) primer, using
methylated C in the first strand synthesis reaction.
Following this first strand reaction, double-stranded cDNA
was blunted, ligated to NotI adapters, digested with
EcoRI, size-selected, and cloned into the NotI and EcoRI
compatible sites of a custom modified MCS of the
pBluescript (KS+) vector. The library was normalized in 2
rounds using conditions adapted from Soares et al., PNAS
(1994) 91: 9228-9232 and Bonaldo et al., Genome Research 6
(1996): 791, except that a significantly longer
reannealing hybridization was used."

ORIGIN

Query Match 28.4%; Score 282.8; DB 13; Length 739;
Best Local Similarity 86.5%; Pred. No. 7.9e-41;
Matches 384; Conservative 0; Mismatches 47; Indels 13; Gaps 6;
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DB 256 GTTCCAGAGGCCCAATGTAGGAGGCTTTTCCACATGCGCAGATGTTGGGAGAGCGA 315
QY 503 TGGAGTCTTAGTATCAGTCATGACAGATGAAGAGGACAGAAAT-----AAATGTTTT 556
DB 316 TGGAAACCTTAGTGACGGTGATGACTGACGACAAAGGTGTAGTAATAGCAAGATGTTGTTT 375
QY 557 ACAAC-TCCTGATTCCTGATGTTTATATATATATATATATATATATATATATATATATAT 615
DB 376 ACACTTCAGGCGTCCCGCATGTTTATATATATATATATATATATATATATATATATAT 435
QY 616 TAAGAGTTTACAG-AAATAAATCTATATTTTGTGAAGGCTAGTGGTATTACTGTAG 674
DB 436 TAAGAGTTTACAGAAACAAATCTATATTTTGTGAAGGCTAGTGGTATTACTGTAG 495
QY 675 ATTTTCAGTAGTTT--CTAAGTCTGTTATTTGTTTAAACAATGCGAGTTTTCACGCTCT 733
DB 496 ATTTTCAGTAGTTTCTTAACTGCTGTTCTGTTTAAACAATGCGAGTTTTCACGCTCT 555


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QY 734 ATGCAATTGTACAAAAAGTTTATAGAAAACTACATGTAAATCTTGATAGCTAAATAAC 793
Db 556 ATGCAATTGTACAAAA--TTATAGAAAACTACATGTAAATCTTGATAGCTGAATAAC 613
QY 794 TTGCATTTCTTTATATAGAACGATTTTGGTTGTTTA--AAATTTATAACAGTTATA 851
Db 614 TTGCATTTCTTTATATAGAACGATTTTGGTTGTTTA--AAATTTATAACAGTTATA 673
QY 852 AAGAAAGATTTATAAGGAAAAAG 875
Db 674 AAGAAAGATCTAACTAAAAAG 697

RESULT 13
LOCUS AI008807/c
DEFINITION EST203258 Normalized rat embryo, Bento Soares Rattus sp. cDNA clone
ACCESSION AI008807
VERSION REWED56 3' end, mRNA sequence.
KEYWORDS Rattus sp.
SOURCE Rattus sp.
ORGANISM Rattus sp.
REFERENCE 1 (bases 1 to 500)
AUTHORS Lee,N.H., Glodek,A., Chandra,I., Mason,T.M., Quackenbush,J.,
Kerlavage,A.R. and Adams,M.D.
TITLE Rat Genome Project: Generation of a Rat EST (RST) Catalog & Rat
JOURNAL Gene Index
COMMENT Unpublished (1998)
Contact: Lee, NH
The Institute for Genomic Research
9712, Medical Center Drive, Rockville, MD 20850, USA
Tel: (301)-838-3529
Fax: (301)-838-0208
Email: nhlee@tigr.org
Seq primer: M13-21.

FEATURES
source
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/organism="Rattus sp."
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/db_xref="taxon:10118"
/clone="REMBD56"
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/clone_lib="Normalized rat embryo, Bento Soares"
/note="Vector: pT7T3Pac; Site_1: EcoRI; Site_2: NotI"

ORIGIN
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Best Local Similarity 95.4%; Pred. No. 1e-40;
Matches 291; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 439 CTTTGTGTTCCAGGACACAATGTAGGAAGCTTTTCCACATGGCAGATGTTGGGCAGA 498
Db 305 CCTAGTTCAAGAGGACACAATGTAGGAAGCTTTTCCACATGGCAGATGTTGGGCAGA 246

QY 499 GCGATGGAGTCCTTATGATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTAC 558
Db 245 GCGATGGAGTCCTTATGATCAGTCATGATGATGAGAGGAGCAGATAAATGTTTAC 186

QY 559 AACTCTGATCCCGCATGGTTTATAATATTCATATACAAAGAGGATTAGACAGTAA 618
Db 185 AACTCTGATCCCGCATGGTTTATAATATTCGTCACAAAGAGGATTAGACAGTAA 126

QY 619 GAGTTTACAGAAATAAATCTATTTTGTGAAGGAGTAGTGATTTACTGTAGATT 678
Db 125 GCGTTTACAGAAATAAATCTATTTTGTGAAGGAGTAGTGATTTACTGTAGATT 66

QY 679 CAGTAGTTTCTAGTCTGTTATTTGTTTAAACAAATGGCAGGTTTACACGCTCTATGCA 738

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Db 65 CAGTAGTTTCTAGTCTGTTATTTGTTTAAACAATGGCAGGTTTACACGCTCTATGCA 6
QY 739 ATTGT 743
Db 5 ATTGT 1

RESULT 14
LOCUS BG795230/c
DEFINITION BG795230 576 bp mRNA linear EST 16-MAY-2001
cDNA_clone UTSW_SM30E11, mRNA sequence.
ACCESSION BG795230
VERSION BG795230.1 GI:14130800
KEYWORDS Mus musculus (house mouse)
SOURCE EST.
ORGANISM Mus musculus
REFERENCE 1 (bases 1 to 576)
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
Gallardo,T.D., Schageman,J.J., Pertsemilidis,A., Garner,H.R.,
Williams,R.S. and Shonet,R.V.
TITLE UT Southwestern Medical Center, Adult Mouse Skeletal Muscle cDNA
JOURNAL Library
COMMENT Unpublished (2001)
Contact: Schageman JJ
Shonet/Garner Labs
University of Texas Southwestern Medical Center
6000 Harry Hines Blvd., NA2.226, Dallas, TX 75390, USA
Tel: 214 648 1674
Email: Jeff.Schageman@UTSouthwestern.edu
cDNA library constructed by UTSW as a component of the Program for
Genomic Applications (PGA) and the Reynolds Heart Disease
Prevention grants for use in cDNA microarray experiments. Sequence
Quality: Sequence ends were trimmed based on percentage of ambigu-
us base calls or N's in windowed segments. Sequencing: First-pass
sequencing; ABI Prism 377 sequencer and analysis software.
Seq primer: M13/pUC Reverse.

FEATURES
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/dev_stage="2 months"
/lab_host="DH5a"
/clone_lib="UTSW Adult Mouse Skeletal Muscle Library"
/note="Vector: pAMP10 (Gibco); Cloned unidirectionally.
Primer: Oligo dt. RNA isolation: cytoplasmic RNA preps
(Manniat); Cloning technique: CUA Cloning (CloneAmp,
Life Technologies); Average insert size: 1.8 Kb;
Insertion site: TAGCTCACTGAATCTGAGTG---. Other
information regarding entire library may be found at
http://pga.swmed.edu/data/libraries/microarray_cdna_librar-
ies.htm."

ORIGIN
Query Match 28.2%; Score 281.2; DB 12; Length 576;
Best Local Similarity 95.8%; Pred. No. 1.7e-40;
Matches 297; Conservative 0; Mismatches 12; Indels 1; Gaps 1;

QY 450 AGGACACAATGTAGGAAGCTTTTCCACATGGCAGATGATTGGGCAGCGATGGATC 509
Db 347 AGGNCNCAATGTAGGAAGCCCTTTTCCACATGGCANATGATTGGGCAGCGATGGATC 288

QY 510 CTTAGTATCAGTCATGACATGAAGAAGGAGCAGATAAATGTTTACAACTCCTGATT 569
Db 287 CTTAGTATCAGTCATGACATGAAGAAGGAGCAGATAAATGTTTACAACTCCTGATT 238

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 11:33:41 ; Search time 2486.92 Seconds
(without alignments)
1500.955 Million cell updates/sec

Title: US-09-966-264D-2

Perfect score: 996

Sequence: 1 GTGGTTGATTGATAGTAA.....GTGTTGATGTTAATTAATT 996

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2470632 seqs, 1873875610 residues

Total number of hits satisfying chosen parameters: 4941264

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : Published applications NA:

- 1: /cgn2_6/ptodata/1/pubpna/US07_PUBCOMB.seq.*
- 2: /cgn2_6/ptodata/1/pubpna/PCT_NEW_PUB.seq.*
- 3: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq.*
- 4: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq.*
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- 18: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	996	100.0	996	US-09-966-264-2	Sequence 2, Appli
2	410.8	41.2	5339	US-10-149-736-40	Sequence 40, Appl
3	410.8	41.2	5417	US-10-149-736-39	Sequence 39, Appl
4	410.8	41.2	5462	US-10-149-736-41	Sequence 41, Appl
5	410.8	41.2	8689	US-10-149-736-42	Sequence 42, Appl
6	410.8	41.2	11443	US-10-149-736-44	Sequence 44, Appl
7	410.8	41.2	12057	US-10-149-736-47	Sequence 47, Appl
8	410.8	41.2	13957	US-09-782-378A-22	Sequence 22, Appl
9	410.8	41.2	13957	US-09-880-107-2284	Sequence 2284, Ap
10	410.8	41.2	13957	US-10-149-736-1	Sequence 1, Appli
11	410.8	41.2	14069	US-10-342-887-434	Sequence 434, App
12	410.8	41.2	14082	US-10-342-887-981	Sequence 981, App
13	410.8	41.2	14082	US-10-341-434-108	Sequence 108, App
14	396.8	39.8	2891	US-10-149-736-38	Sequence 38, Appl
15	374.2	37.6	13815	US-10-149-736-2	Sequence 2, Appli

C 16	151.4	15.2	430	9	US-09-796-692-3505	Sequence 3505, Ap
C 17	151.4	15.2	430	14	US-10-040-862-3505	Sequence 3505, Ap
C 18	151.4	15.2	430	15	US-10-057-475B-3505	Sequence 3505, Ap
C 19	151.4	15.2	430	15	US-10-154-884B-3505	Sequence 3505, Ap
C 20	151.4	15.2	2891	15	US-10-149-736-38	Sequence 38, Appl
C 21	151.4	15.2	8689	15	US-10-149-736-42	Sequence 42, Appl
C 22	151.4	15.2	11443	15	US-10-149-736-44	Sequence 44, Appl
C 23	151.4	15.2	13957	9	US-09-782-378A-22	Sequence 22, Appl
C 24	151.4	15.2	13957	9	US-09-880-107-2284	Sequence 2284, Ap
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C 26	151.4	15.2	14069	12	US-10-342-887-434	Sequence 434, App
C 27	151.4	15.2	14082	12	US-10-342-887-981	Sequence 981, App
C 28	151.4	15.2	14082	15	US-10-341-434-108	Sequence 108, App
C 29	137	13.8	137	9	US-09-966-264-1	Sequence 1, Appli
C 30	101.4	10.2	13815	15	US-10-149-736-2	Sequence 2, Appli
C 31	51.2	5.1	3673778	14	US-10-312-841-1	Sequence 1, Appli
C 32	50.6	5.1	12393	14	US-10-311-455-1236	Sequence 1236, Ap
C 33	49.8	5.0	18154	14	US-10-311-455-228	Sequence 228, App
C 34	49.4	5.0	622	12	US-10-424-593-3380	Sequence 3380, Ap
C 35	48.4	4.9	23683	14	US-10-240-485-176	Sequence 176, App
C 36	48.2	4.8	5546	14	US-10-311-455-364	Sequence 364, App
C 37	48	4.8	8076	16	US-10-257-166-36	Sequence 16, Appl
C 38	47.4	4.8	5413	12	US-10-221-714A-418	Sequence 418, App
C 39	47.4	4.8	3673778	14	US-10-312-841-1	Sequence 1, Appli
C 40	47.4	4.8	3673778	14	US-10-312-841-2	Sequence 1455, Ap
C 41	47.2	4.7	17213	14	US-10-311-455-1455	Sequence 2075, Ap
C 42	47	4.7	4993	14	US-10-311-455-2075	Sequence 57, Appl
C 43	47	4.7	7348	12	US-10-221-714A-57	Sequence 1797, Ap
C 44	47	4.7	8771	14	US-10-311-455-1797	Sequence 1780, Ap
C 45	46.8	4.7	5683	14	US-10-311-455-1780	

ALIGNMENTS

RESULT 1
US-09-966-264-2
; Sequence 2, Application US/09966264
; Patent No. US20020099015A1
; GENERAL INFORMATION:
; APPLICANT: Barber, Elizabeth K
; TITLE OF INVENTION: Gene Expression Control Element DNA
; FILE REFERENCE: 896034605001
; CURRENT APPLICATION NUMBER: US/09/966,264
; CURRENT FILING DATE: 2001-09-28
; PRIOR APPLICATION NUMBER: US 60/237,079
; PRIOR FILING DATE: 2000-09-30
; NUMBER OF SEQ ID NOS: 33
; SOFTWARE: Patent in version 3.1
; SEQ ID NO 2
; LENGTH: 996
; TYPE: DNA
; ORGANISM: human
; FEATURE:
; NAME/KEY: exon
; LOCATION: (1)..(996)
; OTHER INFORMATION:
; NAME/KEY: misc feature
; LOCATION: (710)..(996)
; OTHER INFORMATION: Nucleotides 710-996 are homologous to a portion of human dystroph
; OTHER INFORMATION: in DNA in the region of exon 79 except that nucleotides 860-996 a
; OTHER INFORMATION: re inverted in comparison to the orientation of the same sequence
; OTHER INFORMATION: in the dystrophin DNA
US-09-966-264-2

Query Match 100.0%; Score 996; DB 9; Length 996;
Best Local Similarity 100.0%; Pred. No. 6.9e-224; Indels 0; Gaps 0;
Matches 996; Conservative 0; Mismatches 0;
Qy 1 GTGGTTGATTGATAGTAA.....GTGTTAATTAATCAAGTAGAGAGTAAGTAATCAAT 60
Db 1 GTGGTTGATTGATAGTAA.....GTGTTAATTAATCAAGTAGAGAGTAAGTAATCAAT 60

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QY 61 CAATCACTATAGCAAGGTGAAAGATGATCCCATCATGGAATATCTCTGTCTGAT 120
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QY 241 ACAACACGCTGATGCTTCAATTAAGGCTTCTCTCTCTGGAACCTGCTGAA 300
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QY 421 AAGTCTGTCTTCTTCTCTTCTTCTTCCAGGACACCAATGTAGGAAGCTCTTCCACATG 480
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DB 841 TAACAGTTTAAAGAAAGATTTAATAAGGAAAGAAATAAAGCAATGGAAGTGGTG 900
QY 901 AGCTGTGAATCAGTGTGCAATTTATCAGGAACACCCCAACCAAGTGAAGTGA 960
DB 901 AGCTGTGAATCAGTGTGCAATTTATCAGGAACACCCCAACCAAGTGAAGTGA 960
QY 961 AATAGCATGAGAGCGCTGTTGATCTTAATTAAT 996
DB 961 AATAGCATGAGAGCGCTGTTGATCTTAATTAAT 996

```

```

RESULT 2
US-10-149-736-40
; Sequence 40, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences

```

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; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; PRIOR FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 40
; LENGTH: 5339
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
;
US-10-149-736-40

Query Match 41.2%; Score 410.8; DB 15; Length 5339;
Best Local Similarity 98.3%; Pred. No. 5.7e-86;
Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 450 AGGACACAATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGGAGAGCGATGAGTC 509
DB 4491 AGGACACAATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGGAGAGCGATGAGTC 4550
QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACCTCTGAT 569
DB 4551 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACCTCTGAT 4610
QY 570 CCGCATGTTTATTAATATTCATACAAAGAGGATTAGACAGTAAAGTTTACAAG 629
DB 4611 CCGCATGTTTATTAATATTCATACAAAGAGGATTAGACAGTAAAGTTTACAAG 4670
QY 630 AATAAATCTATATTTTGTGAAGGCTAGTGTATTTATCTAGTATTTTCACTAGTATTTCT 689
DB 4671 AATAAATCTATATTTTGTGAAGGCTAGTGTATTTATCTAGTATTTTCACTAGTATTTCT 4730
QY 690 AAGTCTGTATTTGTTTGTAAACATGGCAGTGTTTTACAGCTGTAGTCAATTTGTCACAAA 749
DB 4731 AAGTCTGTATTTGTTTGTAAACATGGCAGTGTTTTACAGCTGTAGTCAATTTGTCACAAA 4790
QY 750 AAGTTTAAAGAAATACATGTAATAATCTTGTAGCTAAATTAATCTTGTCCATTTCTTTATA 809
DB 4791 AAGTTTAAAGAAATACATGTAATAATCTTGTAGCTAAATTAATCTTGTCCATTTCTTTATA 4850
QY 810 TGAACGCAATTTGGTGTGTTTAAATTTTAAACAGTTTATTAAGAAAGATTTATAAGG 869
DB 4851 TGAACGCAATTTGGTGTGTTTAAATTTTAAACAGTTTATTAAGAAAGATTTATAAGG 4910
QY 870 AA 871
DB 4911 AA 4912

RESULT 3
US-10-149-736-39
; Sequence 39, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; PRIOR FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 39
; LENGTH: 5417

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QY	630	AAATAAATCTATAATTTTGTGAAGGGTAGTGGTATTATTA	CTGTAGATTTCAAGTAGTTTCT	869
Db	6165	AAATAAAATCTATAATTTTGTGAAGGGTAGTGGTATTATTA	CTGTAGATTTCAAGTAGTTTCT	6224
QY	690	AACTCTGTTATTGTTTTGTTTAAACAATGGCAGGTTTTTAC	ACGTCCTATGCAAAATTCACAAA	749
Db	6225	AACTCTGTTATTGTTTTGTTTAAACAATGGCAGGTTTTTAC	ACGTCCTATGCAAAATTCACAAA	6284
QY	750	AACTTATAGAAGAAACATCACTGTAATAATCTTGTATAGCT	TAATAACTTGCCATTCTTTTATA	809
Db	6285	AACTTATAGAAGAAACATCACTGTAATAATCTTGTATAGCT	TAATAACTTGCCATTCTTTTATA	6344
QY	810	TGGAACGCATTTTGGGTTGTTTTAAAAATTTATACAGTT	ATAAGAAAGAAATTTATAAGG	869
Db	6345	TGGAACGCATTTTGGGTTGTTTTAAAAATTTATACAGTT	ATAAGAAAGAAATTTATAAGG	6404
QY	870	AA 871		
Db	6405	AA 6406		

RESULT 6

US-10-149-736-44
; Sequence 44, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; APPLICANT: Harper, Scott Q
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UN-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; CURRENT FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 44
; LENGTH: 11443
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-10-149-736-44

Qy	810	TGGAAGCGCATTTTGGGTTGTTTAAAAATTTATAACAGTTATAAGAAAGAAATTTATAAAGG	869
Db	9099	TGGAAGCGCATTTTGGGTTGTTTAAAAATTTATAACAGTTATAAGAAAGATTGTAAACTA	9158
Qy	870	AA 871	
Db	9159	AA 9160	

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RESULT 7
US-10-149-736-47
; Sequence 47, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; APPLICANT: Harper, Scott Q.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; CURRENT FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 47
; LENGTH: 12057
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-10-149-736-47

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RESULT 8

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US-09-782-378A-22
; Sequence 22, Application US/09782378A
; Patent No. US20020102731A1
; GENERAL INFORMATION:
; APPLICANT: Hearing, Patrick
; APPLICANT: Babou, Wadie
; APPLICANT: Sandaon, Ziv
; APPLICANT: Gnatenko, Dmitri
; TITLE OF INVENTION: Adenoviral Vectors
; FILE REFERENCE: STONYB-04970
; CURRENT APPLICATION NUMBER: US/09/782,378A
; PRIOR FILING DATE: 2001-02-12
; PRIOR APPLICATION NUMBER: 60/237,747
; PRIOR FILING DATE: 2000-10-02
; NUMBER OF SEQ ID NOS: 27
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 22
; LENGTH: 13957
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-782-378A-22

Query Match      41.2%; Score 410.8; DB 9; Length 13957;
Best Local Similarity 98.3%; Pred. No. 9.3e-86;
Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 450 AGGACACAATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGCGAGCGATGGAGTC 509
DB 11253 AGGACACAATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGCGAGCGATGGAGTC 11312

QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 569
DB 11313 CTTAGTATCAGTCATGACAGATGAAGAAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 11372

QY 570 CCCGATGGTCTTATATATATATATATATATATATATATATATATATATATATATAT 629
DB 11373 CCCGATGGTCTTATATATATATATATATATATATATATATATATATATATATATAT 11432

QY 630 AAATAAATCTATATTTTGTGAAGGAGTGTGATTTACTGTAGATTTTCACTAGTTTCT 689
DB 11433 AAATAAATCTATATTTTGTGAAGGAGTGTGATTTACTGTAGATTTTCACTAGTTTCT 11492

QY 690 AAGTCTGTATTTGTTTGTGAAGGAGTGTGATTTACTGTAGATTTTCACTAGTTTCT 749
DB 11493 AAGTCTGTATTTGTTTGTGAAGGAGTGTGATTTACTGTAGATTTTCACTAGTTTCT 11552

QY 750 AAGTTTAAAGAAACTACATGTAAATCTTGTAGTAAATCTTGTAGTAAATCTTGTAGTAA 809
DB 11553 AAGTTTAAAGAAACTACATGTAAATCTTGTAGTAAATCTTGTAGTAAATCTTGTAGTAA 11612

QY 810 TGGACGCAATTTGGTGTGTTTAAATTTTAAATTTTAAATTTTAAATTTTAAATTTTAA 869
DB 11613 TGGACGCAATTTGGTGTGTTTAAATTTTAAATTTTAAATTTTAAATTTTAAATTTTAA 11672

QY 870 AA 871
DB 11673 AA 11674

RESULT 9
US-09-880-107-2284
; Sequence 2284, Application US/09880107
; Patent No. US20020142981A1
; GENERAL INFORMATION:
; APPLICANT: Horne, Daryl T.
; APPLICANT: Vockley, Joseph G.
; APPLICANT: Scherf, Uwe
; APPLICANT: Gene Logic, Inc.
; TITLE OF INVENTION: Gene Expression Profiles in Liver Cancer
; FILE REFERENCE: 44921-5028-WO
; CURRENT APPLICATION NUMBER: US/09/880,107
; CURRENT FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/211,379

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; PRIOR FILING DATE: 2000-06-14
; PRIOR APPLICATION NUMBER: US 60/237,054
; PRIOR FILING DATE: 2000-10-02
; NUMBER OF SEQ ID NOS: 3950
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2284
; LENGTH: 13957
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: Genbank Accession No. US20020142981A1 M18533
US-09-880-107-2284

Query Match      41.2%; Score 410.8; DB 9; Length 13957;
Best Local Similarity 98.3%; Pred. No. 9.3e-86;
Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 450 AGGACACAATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGCGAGCGATGGAGTC 509
DB 11253 AGGACACAATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGCGAGCGATGGAGTC 11312

QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 569
DB 11313 CTTAGTATCAGTCATGACAGATGAAGAAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG 11372

QY 570 CCCGATGGTCTTATATATATATATATATATATATATATATATATATATATATATAT 629
DB 11373 CCCGATGGTCTTATATATATATATATATATATATATATATATATATATATATATAT 11432

QY 630 AAATAAATCTATATTTTGTGAAGGAGTGTGATTTACTGTAGATTTTCACTAGTTTCT 689
DB 11433 AAATAAATCTATATTTTGTGAAGGAGTGTGATTTACTGTAGATTTTCACTAGTTTCT 11492

QY 690 AAGTCTGTATTTGTTTGTGAAGGAGTGTGATTTACTGTAGATTTTCACTAGTTTCT 749
DB 11493 AAGTCTGTATTTGTTTGTGAAGGAGTGTGATTTACTGTAGATTTTCACTAGTTTCT 11552

QY 750 AAGTTTAAAGAAACTACATGTAAATCTTGTAGTAAATCTTGTAGTAAATCTTGTAGTAA 809
DB 11553 AAGTTTAAAGAAACTACATGTAAATCTTGTAGTAAATCTTGTAGTAAATCTTGTAGTAA 11612

QY 810 TGGACGCAATTTGGTGTGTTTAAATTTTAAATTTTAAATTTTAAATTTTAAATTTTAA 869
DB 11613 TGGACGCAATTTGGTGTGTTTAAATTTTAAATTTTAAATTTTAAATTTTAAATTTTAA 11672

QY 870 AA 871
DB 11673 AA 11674

RESULT 10
US-10-149-736-1
; Sequence 1, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; APPLICANT: Harper, Scott Q.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; CURRENT FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
; LENGTH: 13957
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-149-736-1

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QY 570 CCGCATGGTTTTTATAATTATTCATACAAAGAGGATTAGACAGTAGTAAAGTTTACAA 629
Db 11498 CCGCATGGTTTTTATAATTATTCATACAAAGAGGATTAGACAGTAGTAAAGTTTACAA 11557
QY 630 AAATAAATCTATATTTTGTGAAGGAGTGTATTTACTGTAGATTTCAAGTATTTCT 689
Db 11558 AAATAAATCTATATTTTGTGAAGGAGTGTATTTACTGTAGATTTCAAGTATTTCT 11617
QY 690 AAGTCTGTTATTTGTTTAAACAATGGCAGGTTTTTACACGCTCTATGCAATTTGACAAAA 749
Db 11618 AAGTCTGTTATTTGTTTAAACAATGGCAGGTTTTTACACGCTCTATGCAATTTGACAAAA 11677
QY 750 AAGTTATAGAAACTACATGTAATAATCTGTAGCTAAATAACTTGCATTTCTTTATA 809
Db 11678 AAGTTATAGAAACTACATGTAATAATCTGTAGCTAAATAACTTGCATTTCTTTATA 11737
QY 810 TGGAAACGCAATTTGGTGTGTTTAAAAATTTATAACAGTTATAAGAAAGAAATTTAAAGG 869
Db 11738 TGGAAACGCAATTTGGTGTGTTTAAAAATTTATAACAGTTATAAGAAAGAAATTTAAAGG 11797
QY 870 AA 871
Db 11798 AA 11799
RESULT 13
US-10-341-434-108
; Sequence 108, Application US/10341434
; Publication No. US20030215835A1
; GENERAL INFORMATION:
; APPLICANT: Origene Technologies
; TITLE OF INVENTION: Differentially Regulated Prostate Cancer Genes
; FILE REFERENCE: 9U 204 205 R1
; CURRENT APPLICATION NUMBER: US/10/341,434
; CURRENT FILING DATE: 2003-07-18
; PRIOR APPLICATION NUMBER: US 60/348,164
; PRIOR FILING DATE: 2002-01-15
; PRIOR APPLICATION NUMBER: US 60/348,119
; PRIOR FILING DATE: 2002-01-15
; NUMBER OF SEQ ID NOS: 238
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 108
; LENGTH: 14082
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (703)..(11388)
; OTHER INFORMATION:
US-10-341-434-108
Query Match 41.2%; Score 410.8; DB 15; Length 14082;
Best Local Similarity 98.3%; Pred. No. 9.4e-86;
Matches 415; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
QY 450 AGACACAATGTAGGAAGTCTTTCCACATGCGAGATGATTTGGCAGAGCGATGGAGTC 509
Db 11378 AGACACAATGTAGGAAGTCTTTCCACATGCGAGATGATTTGGCAGAGCGATGGAGTC 11437
QY 510 CTTAGTATCAGTCATGACAGATGAGAGGAGCAGATAAATGTTTCAACTCCTCGATT 569
Db 11438 CTTAGTATCAGTCATGACAGATGAGAGGAGCAGATAAATGTTTCAACTCCTCGATT 11497
QY 570 CCGCATGGTTTTTATAATTATTCATACAAAGAGGATTAGACAGTAGTAAAGTTTACAA 629
Db 11498 CCGCATGGTTTTTATAATTATTCATACAAAGAGGATTAGACAGTAGTAAAGTTTACAA 11557
QY 630 AAATAAATCTATATTTTGTGAAGGAGTGTATTTACTGTAGATTTCAAGTATTTCT 689
Db 11558 AAATAAATCTATATTTTGTGAAGGAGTGTATTTACTGTAGATTTCAAGTATTTCT 11617
QY 690 AAGTCTGTTATTTGTTTAAACAATGGCAGGTTTTTACACGCTCTATGCAATTTGACAAAA 749

Db 11618 AAGTCTGTTATTTGTTTAAACAATGGCAGGTTTTTACACGCTCTATGCAATTTGACAAA 11677
QY 750 AAGTTATAGAAACTACATGTAATAATCTGTAGCTAAATAACTTGCATTTCTTTATA 809
Db 11678 AAGTTATAGAAACTACATGTAATAATCTGTAGCTAAATAACTTGCATTTCTTTATA 11737
QY 810 TGGAAACGCAATTTGGGTTGTTTAAAAATTTATAACAGTTATAAGAAAGAAATTTAAAGG 869
Db 11738 TGGAAACGCAATTTGGGTTGTTTAAAAATTTATAACAGTTATAAGAAAGAAATTTAAAGG 11797
QY 870 AA 871
Db 11798 AA 11799
RESULT 14
US-10-149-736-38
; Sequence 38, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; CURRENT FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 38
; LENGTH: 2691
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-149-736-38
Query Match 39.8%; Score 396.8; DB 15; Length 2691;
Best Local Similarity 98.3%; Pred. No. 7.8e-83;
Matches 401; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
QY 464 GAAGTCTTTTCCATGGCAGATGATTTGGCAGAGCGATGGAGTCTTAGTATCAGTCA 523
Db 1 GAAGTCTTTTCCATGGCAGATGATTTGGCAGAGCGATGGAGTCTTAGTATCAGTCA 60
QY 524 TGACAGATGAGAGGAGCAGATAAATCTTTTACAACTCCTGATTCGCCATGGTTTTT 583
Db 61 TGACAGATGAGAGGAGCAGATAAATCTTTTACAACTCCTGATTCGCCATGGTTTTT 120
QY 584 ATAATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAGAAATAAATCTATAT 643
Db 121 ATAATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAGAAATAAATCTATAT 180
QY 644 TTTTGTGAAGGAGTGGTATTTACTAGTAGATTTTCAAGTATTTCTAAGTCTGTTATGT 703
Db 181 TTTTGTGAAGGAGTGGTATTTACTAGTAGATTTTCAAGTATTTCTAAGTCTGTTATGT 240
QY 704 TTTTGTAAACATGGCAGGTTTTTACAGTCTATGCAATTTGACAAAAAGTTATAAGAAA 763
Db 241 TTTTGTAAACATGGCAGGTTTTTACAGTCTATGCAATTTGACAAAAAGTTATAAGAAA 300
QY 764 CTACATGTAATAATCTTTGATAGCTAAATACTTGCATTTCTTTTATATGGAACCCATTTG 823
Db 301 CTACATGTAATAATCTTTGATAGCTAAATACTTGCATTTCTTTTATATGGAACCCATTTG 360
QY 824 GGTGTTTAAAAATTTATAACAGTTATAAGAAAGAAATTTATAAGAA 871
Db 361 GGTGTTTAAAAATTTATAACAGTTATAAGAAAGAAATTTATAAGAA 408
RESULT 15
US-10-149-736-2

; Sequence 2, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; CURRENT FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 13815
; TYPE: DNA
; ORGANISM: Mus musculus
US-10-149-736-2

Query Match 37.6%; Score 374.2; DB 15; Length 13815;
Best Local Similarity 96.0%; Pred. No. 3.7e-77;
Matches 406; Conservative 0; Mismatches 13; Indels 4; Gaps 2;

QY 450 AGGACACAATGTAGGAAGTCCTTTCCACATGCGCAGATGATTTGGCGAGCGATGGAGTC 509
DB 11244 AGGACACAATGTAGGAAGCCCTTTCCACATGCGCAGATGATTTGGCGAGCGATGGAGTC 11303

QY 510 CTTAGTATCAGTCATGACAGATGAGAGAGGAGCAGATAAATGTTTACAACTCCTGATT 569
DB 11304 CTTAGTTCAGTCATGACAGATGAGAGAGGAGCAGATAAATGTTTACAACTCCTGATT 11363

QY 570 CCCGCATGCTTTTATATATTTATATATTTATATATTTATATATTTATATATTTATATATTT 629
DB 11364 CCCGCATGCTTTTATATATTTATATATTTATATATTTATATATTTATATATTTATATATTT 11423

QY 630 AAAT-AAATCTATATTTTGTGAAGGAGTAGTGGTATTTATCTAGTATGATTTTCAGTAGTTTC 688
DB 11424 AAATAAATCTATATTTTGTGAAGGAGTAGTGGTATTTATCTAGTATGATTTTCAGTAGTTTC 11483

QY 689 TAAGTCTGTTATTTGTTTAACTAATGCGAGGTTTACACGTCCTATGCAATTTGTACAAA 748
DB 11484 TAAGTCTGTTATTTGTTTAACTAATGCGAGGTTTACACGTCCTATGCAATTTGTACAAA 11543

QY 749 AAAGTTATAAGAAACTACATGTAATAATCTTGATAGCTAAATAACTTCCCATTTCTTTAT 808
DB 11544 AAAGTTAAAAGAAA---ACATGTAATAATCTTGATAGCTAAATAACTTCCCATTTCTTTAT 11600

QY 809 ATGGAACGCATTTTGGGTTGTTTAAATAATTTATATACAGTTTATAAGAAAGAAATTATAAG 868
DB 11601 ATGGAACGCATTTTGGGTTGTTTAAATAATTTATATACAGTTTATAAGAAAGAAATTATAAG 11660

QY 869 GAA 871
DB 11661 AAA 11663

Search completed: April 6, 2004, 15:01:46
Job time : 2493.92 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 14:12:12 ; Search time 3672.8 Seconds

(without alignments)
8098.097 Million cell updates/sec

Title: US-09-966-264D-2

Perfect score: 996

Sequence: 1 ggggttgatgtagtaaa.....gtgttgatgtagtaatt 996

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 27513289 seqs, 14931090276 residues

Word size : 0

Total number of hits satisfying chosen parameters: 55026578

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

1: em_estba.*

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4: em_estnu.*

5: em_estov.*

6: em_estpl.*

7: em_estro.*

8: em_htc.*

9: gb_est1.*

10: gb_est2.*

11: gb_hcc.*

12: gb_est3.*

13: gb_est4.*

14: gb_est5.*

15: em_estfun.*

16: em_estom.*

17: em_gss_hum.*

18: em_gss_inv.*

19: em_gss_pln.*

20: em_gss_vrt.*

21: em_gss_fun.*

22: em_gss_nam.*

23: em_gss_mus.*

24: em_gss_pro.*

25: em_gss_rtd.*

26: em_gss_phg.*

27: em_gss_vrl.*

28: gb_gsl1.*

29: gb_gss2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
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2	342	34.3	629	9	AL712236 602669567
3	342	34.3	629	13	AL712236 602669567
4	238	23.9	999	13	BX419180 602669567

5	231	23.2	739	9	AU138648
6	221	22.2	308	14	N75050
7	209	21.0	275	10	AW604756
8	209	21.0	277	10	AW861957
9	209	21.0	284	10	AW858457
10	208	20.9	888	14	CF552038
11	202	20.3	525	28	AQ679243
12	194	19.5	271	10	AW858578
13	183	18.4	619	10	AW771158
14	161	16.2	309	14	H27701
15	148	14.9	556	28	AQ504173
16	145	14.7	396	14	H89576
17	145	14.7	455	9	AA427831
18	146	14.7	482	14	CB045405
19	146	14.7	473	14	CA389721
20	146	14.7	489	28	AQ015185
21	146	14.7	647	10	AW664684
22	146	14.7	696	10	AW950480
23	146	14.7	710	9	AV725574
24	146	14.7	727	12	BG567176
25	146	14.7	740	14	CD356811
26	146	14.7	745	14	CD357556
27	145	14.7	1121	12	BM546012
28	145	14.6	743	14	CD110642
29	141	14.2	373	10	AW580404
30	129	13.0	939	13	BO718541
31	120	12.0	352	10	AW385154
32	118	11.8	501	13	HU619605
33	112	11.2	862	14	CB962272
34	111	11.1	353	10	AW580423
35	109	10.9	183	10	AW585837
36	97	9.7	911	12	BI752714
37	94	9.4	439	14	CB750007
38	94	9.4	924	13	BQ327942
39	94	9.4	4437	11	AK036936
40	91	9.1	492	14	CB725360
41	91	9.1	571	14	CA893354
42	90	9.0	375	10	AW607064
43	88	8.8	484	10	BF655326
44	87	8.7	531	12	BI281078
45	87	8.7	576	12	BG795230

ALIGNMENTS

RESULT 1
BG706268
LOCUS 602669567F1 NIH_MGC_96 Homo sapiens CDNA clone IMAGE:4792313 5',
DEFINITION 844 bp mRNA linear EST 07-MAY-2001
mRNA sequence.
ACCESSION BG706268
VERSION BG706268.1 GI:13981445
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 844)
AUTHORS NIH-MGC <http://mgs.nci.nih.gov/>
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabs@mail.nih.gov
Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
CDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki Toshiyuki and Piero Carninci (RIKEN)
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: <http://image.llnl.gov>
Plate: LLM10670 row: e column: 18

EST (Poustka, A., Albert, R., Moosmayer, P., Schupp, I.,
Wellenreuther, R., et al.)
Unpublished (2003)
Contact: MIPS

INGOLSTAEDTER Landstr. 1, D-85764 Neuherberg, Germany
This is the 5' sequence of the clone insert
Cloned from S. Wiemann, Molecular Genome Analysis, German Cancer
Research Center (DKFZ); Email: s.wiemann@dkfz-heidelberg.de;
sequenced by DKFZ (German Cancer Research Center,
Heidelberg/Germany) within the cDNA sequencing consortium of the
German Genome Project.
No sl sequence available.
This clone (DKFZ686J2487) is available at the RZPD in Berlin.
Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059
Berlin-Charlottenburg, GERMANY; Email: clone@rzd.de.
Location/Qualifiers
1. 629
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="DKFZ686J2487"
/dev_stage="adult"
/lab_host="DH10B"
/clone_lib="686 (synonym: hlcc3)"
/note="Vector: pTriplex2, Site_1: SfiIA; Site_2: SfiIB;
cDNA-collection"

ORIGIN
Query Match 34.3%; Score 342; DB 13; Length 629;
Best Local Similarity 100.0%; Pred. No. 3e-156;
Matches 342; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 518 CAGTCATCACAGATGACGAGGCGAGTAATGTTTACAACTCTGATCCCGCATG 577
Db 1 CAGTCATCACAGATGACGAGGCGAGTAATGTTTACAACTCTGATCCCGCATG 60

QY 578 GTTTTATATATTCATACACAAAGAGGATTAGACAGTACAGTATTACAGAAATAAAT 637
Db 61 GTTTTATATATTCATACACAAAGAGGATTAGACAGTACAGTATTACAGAAATAAAT 120

QY 638 CTATATTTTGTGAGGAGTGGTATTATATCTAGATTTCAGTAGTTCTTAAGTCTGT 697
Db 121 CTATATTTTGTGAGGAGTGGTATTATATCTAGATTTCAGTAGTTCTTAAGTCTGT 180

QY 698 TATTGTTTGTAACTAGCGAGTTTACACGCTCTATGCAATGTACAAAAGTTTATA 757
Db 181 TATTGTTTGTAACTAGCGAGTTTACACGCTCTATGCAATGTACAAAAGTTTATA 240

QY 758 AGAAACTACATGTAATAATCTGTAGCTAAATACTGGCCATTTCTTTATATGGAACG 817
Db 241 AGAAACTACATGTAATAATCTGTAGCTAAATACTGGCCATTTCTTTATATGGAACG 300

QY 818 ATTTGGGTTGTTTAAATTTATAACAGTATTATAAGAAAGA 859
Db 301 ATTTGGGTTGTTTAAATTTATAACAGTATTATAAGAAAGA 342

RESULT 4
BX419180
LOCUS
DEFINITION
BX419180 Homo sapiens FETAL BRAIN Homo sapiens cDNA clone
CS0DF014YA17 5-PRIME, mRNA sequence.

ACCESSION
BX419180
VERSION
BX419180.1 GI:30637926
KEYWORDS
EST.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 999)
REFERENCE
Li, W.B., Gruber, C., Jesse, J., and Polayes, D.
Full-length cDNA libraries and normalization

Unpublished (2001)
Contact: Genoscope
Genoscope - Centre National de Sequencage
BP 191 91006 EVRY cedex - France
Email: seqref@genoscope.cns.fr, Web: www.genoscope.cns.fr
Library was constructed by Life Technologies, a division of
Invitrogen. This sequence belongs to sequence cluster 10533.r For
more information about this cluster, see
http://www.genoscope.cns.fr/
cgi-bin/cluster.cgi?seq=CS0DF014AA090P1&cluster=10533.r. Contact:
Peng Liang Email: fliang@lifetech.com URL:
http://fulllength.invitrogen.com/ Invitrogen Corporation 1600
Faraday Avenue Genoscope sequence ID: CS0DF014AA090P1.
Location/Qualifiers
1. 999
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="CS0DF014YA17"
/tissue_type="FETAL BRAIN"
/dev_stage="fetal"
/clone_lib="Homo sapiens FETAL BRAIN"
/note="Organ: Brain; Vector: pCMVSPORT 6; 1st strand cDNA
was primed with a NotI-oligo(dT) primer. Five prime end
enriched double-strand cDNA was digested with NotI and
cloned into the NotI and EcoRV sites of the pCMVSPORT 6
vector. Library was not normalized."

ORIGIN
Query Match 23.9%; Score 238; DB 13; Length 999;
Best Local Similarity 100.0%; Pred. No. 2.4e-105;
Matches 238; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 622 TTTCAGAAATAAATCTATATTTTGTGAGGAGTAGTGTATTATATCTAGATTTCAG 681
Db 69 TTTCAGAAATAAATCTATATTTTGTGAGGAGTAGTGTATTATATCTAGATTTCAG 128

QY 682 TAGTTTCTAAGTCTGTATTGTTTGTAACTATGCAATGTCAGCTTACACGCTCTATGCAATT 741
Db 129 TAGTTTCTAAGTCTGTATTGTTTGTAACTATGCAATGTCAGCTTACACGCTCTATGCAATT 188

QY 742 GTACAAAAAGTTTATAAGAAAACTACATGTAATAAATCTTGATAGCTAAATAAATTCGCAATT 801
Db 189 GTACAAAAAGTTTATAAGAAAACTACATGTAATAAATCTTGATAGCTAAATAAATTCGCAATT 248

QY 802 TCTTTATATGGAACGCAATTTGGTGTGTTTAAATTTTATACAGTTTATAAGAAAGA 859
Db 249 TCTTTATATGGAACGCAATTTGGTGTGTTTAAATTTTATACAGTTTATAAGAAAGA 306

RESULT 5
AU138648
LOCUS
DEFINITION
AU138648 PLACE1 Homo sapiens cDNA clone PLACE1009015 5', mRNA
sequence.

ACCESSION
AU138648
VERSION
AU138648.1 GI:11000169
KEYWORDS
EST.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 739)
REFERENCE
Ota, T., Nishikawa, T., Suzuki, Y., Ishii, S., Saito, K., Kawai, Y.,
Yamamoto, J., Wakamatsu, A., Nakamura, Y., Nagai, T., Sugano, S. and
Isogai, T.
HRI human cDNA project
Unpublished (2000)
Contact: Takao Isogai
Genomics Laboratory
Helix Research Institute
1532-3 Yana, Kisarazu, Chiba 292-0812, Japan
Tel: 81-438-52-3975

Fax: 81-438-52-3986
 Email: genomics@ri.co.jp
 RRI human cDNA project; 5'- & 3'-end one pass sequencing: Helix
 Research Institute; cDNA library construction: Department of
 Virology, Institute of Medical Science, University of Tokyo, and
 Helix Research Institute.
 Location/Qualifiers
 1. 739
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="PLACE1009015"
 /tissue_type="placenta"
 /clone_lib="PLACE1"
 /note="Vector: pME18SFL3"

Query Match 23.2%; Score 231; DB 9; Length 739;
 Best Local Similarity 100.0%; Pred. No. 6.8e-102;
 Matches 231; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 629 GAAATAAATCTATATTTTGTGAAGGGTAGTGGTATTATACGTAGATTTCAGTAGTTTC 688
 Db 38 GAAATAAATCTATATTTTGTGAAGGGTAGTGGTATTATACGTAGATTTCAGTAGTTTC 97
 QY 689 TAAGTCGTATGTTTGTGAAGGGTAGTGGTATTATACGTAGATTTCAGTAGTTTC 748
 Db 98 TAAGTCGTATGTTTGTGAAGGGTAGTGGTATTATACGTAGATTTCAGTAGTTTC 157
 QY 749 AAGATTATAGAACTACATGTTTGAAGGGTAGTGGTATTATACGTAGATTTCAGTAGTTTC 808
 Db 158 AAGATTATAGAACTACATGTTTGAAGGGTAGTGGTATTATACGTAGATTTCAGTAGTTTC 217
 QY 809 ATGGAACGCGATTTGGGTTTGAAGGGTAGTGGTATTATACGTAGATTTCAGTAGTTTC 859
 Db 218 ATGGAACGCGATTTGGGTTTGAAGGGTAGTGGTATTATACGTAGATTTCAGTAGTTTC 268

RESULT 6
 N75050/c
 LOCUS
 DEFINITION
 IMAGE:299724 3' similar to gb:M18533 DYSTROPHIN (HUMAN);, mRNA
 sequence.
 N75050
 VERSION
 KEYWORDS
 EST.
 SOURCE
 Homo sapiens (human)
 ORGANISM
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 Hillier, L., Clark, N., Duboue, T., Elliston, K., Hawkins, M.,
 Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M.,
 Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F.,
 Trevaskis, E., Waterston, R., Williamson, A., Wohldmann, P. and
 Wilson, R.
 The WashU-Merck EST Project
 Unpublished (1995)
 CONTACT: Wilson RK
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu
 This clone is available royalty-free through LNL; contact the
 IMAGE Consortium (info@image.lnl.gov) for further information.
 Seq primer: ml3 -40 forward
 High quality sequence stop: 227.
 Location/Qualifiers
 1. 308
 /organism="Homo sapiens"
 /mol_type="mRNA"

FEATURES
 source

/db_xref="GDB:1244648"
 /db_xref="taxon:9606"
 /clone="IMAGE:299724"
 /dev_stage="19 weeks"
 /lab_hosts="DH10B (ampicillin resistant)"
 /clone_lib="Soares fetal_lung_NBHL19W"
 /notes="Organ: lung; Vector: p773D (Pharmacia) with a
 modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st
 strand cDNA was primed with a Not I - cligo(dT) primer
 [5'-TGTTACCAATCTGAAGTGGGCGGCGCAATTTTCTTTTCTTTT-3'],
 double-stranded cDNA was size selected, ligated to Eco RI
 adapters (Pharmacia), digested with Not I and cloned into
 the Not I and Eco RI sites of a modified p773 vector
 (Pharmacia). Library went through one round of
 normalization to a Cot = 5. Library constructed by Bento
 Soares and M. Fatima Bonaldo. This library was constructed
 from the same fetus as the fetal heart library, Soares
 fetal heart NBHL19W."

Query Match 22.2%; Score 221; DB 14; Length 308;
 Best Local Similarity 100.0%; Pred. No. 6.2e-97;
 Matches 221; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 639 TATATTTTCTGAAGGGTAGTGGTATTATACGTAGATTTCAGTAGTTTCAGTAGTTTC 698
 Db 230 TATATTTTCTGAAGGGTAGTGGTATTATACGTAGATTTCAGTAGTTTCAGTAGTTTC 171
 QY 699 ATTGTTTGTAACTAATGCGAGGTTTACACGCTCTATGCAATTTGTAACAAAAGTTTATA 758
 Db 170 ATTGTTTGTAACTAATGCGAGGTTTACACGCTCTATGCAATTTGTAACAAAAGTTTATA 111
 QY 759 GAAACTACATGTAATAATCTTGTAGTAACTAACTTGCATTTCCTTATATGGAACGCA 818
 Db 110 GAAACTACATGTAATAATCTTGTAGTAACTAACTTGCATTTCCTTATATGGAACGCA 51
 QY 819 TTTTGGTGTCTTTAAAAATTTTATAACAGTTTATAAAGAAAGA 859
 Db 50 TTTTGGTGTCTTTAAAAATTTTATAACAGTTTATAAAGAAAGA 10

RESULT 7
 AW604756
 LOCUS
 DEFINITION
 IMAGE:299724 3' similar to gb:M18533 DYSTROPHIN (HUMAN);, mRNA
 sequence.
 N75050
 VERSION
 KEYWORDS
 EST.
 SOURCE
 Homo sapiens (human)
 ORGANISM
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 Hillier, L., Clark, N., Duboue, T., Elliston, K., Hawkins, M.,
 Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M.,
 Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F.,
 Trevaskis, E., Waterston, R., Williamson, A., Wohldmann, P. and
 Wilson, R.
 The WashU-Merck EST Project
 Unpublished (1999)
 CONTACT: Simpson A.J.G.
 Laboratory of Cancer Genetics
 Ludwig Institute for Cancer Research
 Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
 Brazil
 Tel: +55-11-2704922
 Fax: +55-11-2707001
 Email: asimpson@ludwig.org.br
 This sequence was derived from the FAPESP/LICR Human Cancer Genome
 Project. This entry can be seen in the following URL
 (http://www.ludwig.org.br/scripts/gethtml2.pl?tl=CM0&t2=CM0-CT0341-
 260100-160-a10&t3=2000-01-36&t4=1)
 Seq primer: puc 18 forward
 High quality sequence start: 10
 High quality sequence stop: 275.
 Location/Qualifiers
 1. 275

FEATURES
 source

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/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_lib="CT0341"
/note="Organ: colon; Vector: puc18; Site 1: SmaI; Site 2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

ORIGIN
Query Match      21.0%; Score 209; DB 10; Length 275;
Best Local Similarity 100.0%; Pred. No. 4.8e-91;
Matches 209; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 450 AGGACACAATGAGGAAGTCTTTCCACATGGCAGATGTTGGCAGAGCGATGGAGTC 509
DB 55 AGGACACAATGAGGAAGTCTTTCCACATGGCAGATGTTGGCAGAGCGATGGAGTC 114
QY 510 CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTACAACTCCTGATT 569
DB 115 CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTACAACTCCTGATT 174
QY 570 CCCGATGCTTTTATAATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAAG 629
DB 175 CCCGATGCTTTTATAATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAAG 234
QY 630 AAATAAATCTATATTTTGTGAAGGGTAG 658
DB 235 AAATAAATCTATATTTTGTGAAGGGTAG 263

RESULT 8
AW861957
LOCUS      CM3-CT0341-190400-152-c06 CT0341 Homo sapiens cDNA, mRNA sequence.
DEFINITION
ACCESSION  AW861957
VERSION     AW861957.1 GI:7957650
KEYWORDS   EST.
SOURCE      Homo sapiens (human)
ORGANISM    Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE   1 (bases 1 to 277)
AUTHORS     Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,
            Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F.,
            Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H.,
            Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V.,
            O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and
            Simpson,A.J.
            Shotgun sequencing of the human transcriptome with ORF expressed
            sequence tags
            Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
JOURNAL     10737800
MEDLINE     20202663
PUBMED      10737800
COMMENT     Contact: Simpson A.J.G.
            Laboratory of Cancer Genetics
            Ludwig Institute for Cancer Research
            Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
            Brazil
            Tel: +55-11-2704922
            Fax: +55-11-2707001
            Email: asimpson@ludwig.org.br
            This sequence was derived from the FAPESP/LICR Human Cancer Genome
            Project. This entry can be seen in the following URL
            (http://www.ludwig.org.br/scripts/gethtml2.pl?tl=st2-CM3-CT0341-190
            400-152-c06&t3=2000-04-19&t4=1)
            Seq primer: puc 18 forward
            High quality sequence start: 12
            High quality sequence stop: 277.

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FEATURES             Location/Qualifiers
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     organism="Homo sapiens"
     mol_type="mRNA"
     db_xref="taxon:9606"
     dev_stage="Adult"
     clone_lib="CT0341"
     note="Organ: colon; Vector: puc18; Site 1: SmaI; Site 2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."

ORIGIN
Query Match      21.0%; Score 209; DB 10; Length 277;
Best Local Similarity 100.0%; Pred. No. 4.8e-91;
Matches 209; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 450 AGGACACAATGAGGAAGTCTTTCCACATGGCAGATGTTGGCAGAGCGATGGAGTC 509
DB 57 AGGACACAATGAGGAAGTCTTTCCACATGGCAGATGTTGGCAGAGCGATGGAGTC 116
QY 510 CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTACAACTCCTGATT 569
DB 117 CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTACAACTCCTGATT 176
QY 570 CCCGATGCTTTTATAATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAAG 629
DB 177 CCCGATGCTTTTATAATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAAG 236
QY 630 AAATAAATCTATATTTTGTGAAGGGTAG 658
DB 237 AAATAAATCTATATTTTGTGAAGGGTAG 265

RESULT 9
AW858457
LOCUS      CM3-CT0341-170200-093-b02 CT0341 Homo sapiens cDNA, mRNA sequence.
DEFINITION
ACCESSION  AW858457
VERSION     AW858457.1 GI:7954150
KEYWORDS   EST.
SOURCE      Homo sapiens (human)
ORGANISM    Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE   1 (bases 1 to 284)
AUTHORS     Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,
            Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F.,
            Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H.,
            Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V.,
            O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and
            Simpson,A.J.
            Shotgun sequencing of the human transcriptome with ORF expressed
            sequence tags
            Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
JOURNAL     20202663
MEDLINE     10737800
PUBMED      10737800
COMMENT     Contact: Simpson A.J.G.
            Laboratory of Cancer Genetics
            Ludwig Institute for Cancer Research
            Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,
            Brazil
            Tel: +55-11-2704922
            Fax: +55-11-2707001
            Email: asimpson@ludwig.org.br
            This sequence was derived from the FAPESP/LICR Human Cancer Genome
            Project. This entry can be seen in the following URL
            (http://www.ludwig.org.br/scripts/gethtml2.pl?tl=st2-CM3-CT0341-170
            200-093-b02&t3=2000-02-17&t4=1)
            Seq primer: puc 18 forward

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ORIGIN

Query Match 14.9%; Score 148; DB 28; Length 556;
 Best Local Similarity 100.0%; Pred. No. 3.2e-61;
 Matches 148; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy	1	GTGGTTTGATTGATAGTAAATAAATAATGTTGGTTTAAATACAAAGTCAGAGAGTAAGTAATCAAT	60
Db	153	GTGGTTTGATTGATAGTAAATAAATAATGTTGGTTTAAATACAAAGTCAGAGAGTAAGTAATCAAT	94
Qy	61	CAATCACTCATAGCCAAAGTGGAAAAAGATGTATCCCATCATGGAATATTCCTGTTCGTGAT	120
Db	93	CAATCACTCATAGCCAAAGTGGAAAAAGATGTATCCCATCATGGAATATTCCTGTTCGTGAT	34
Qy	121	AGAAATCTTGCTTATCTATGGAATTC	148
Db	33	AGAAATCTTGCTTATCTATGGAATTC	6

Search completed: April 6, 2004, 17:40:09
 Job time : 3674.8 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 13:02:17 ; Search time 4176.52 Seconds
(without alignments)
10336.278 Million cell updates/sec

Title: US-09-966-264D-2
Perfect score: 996
Sequence: 1 gtgtttgatgatgataaa.....gtgtttgatgatttaatt 996

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 3470272 seqs, 21671516995 residues

Word size : 0
Total number of hits satisfying chosen parameters: 6940544

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : GenEmbl.*

- 1: gb.ba.*
- 2: gb.htg.*
- 3: gb.in.*
- 4: gb.om.*
- 5: gb.ov.*
- 6: gb.pat.*
- 7: gb.ph.*
- 8: gb.pl.*
- 9: gb.pr.*
- 10: gb.ro.*
- 11: gb.sts.*
- 12: gb.sv.*
- 13: gb.un.*
- 14: gb.vi.*
- 15: em.ba.*
- 16: em.fun.*
- 17: em.hum.*
- 18: em.in.*
- 19: em.mu.*
- 20: em.om.*
- 21: em.or.*
- 22: em.ov.*
- 23: em.pat.*
- 24: em.ph.*
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- 28: em.un.*
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- 31: em.htg.inv.*
- 32: em.htg.other.*
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- 34: em.htg.pln.*
- 35: em.htg.rod.*
- 36: em.htg.mam.*
- 37: em.htg.vrt.*
- 38: em.sv.*
- 39: em.htgo.hum.*
- 40: em.htgo.mus.*
- 41: em.htgo.other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
C 1	859	86.2	98056	9	AC006061	AC006061 Homo sapi
C 2	808	81.1	21220	2	AC023414	AC023414 Homo sapi
C 3	410	41.2	2463	9	AX817316	AX817316 Sequence
C 4	410	41.2	4658	9	BC028720	BC028720 Homo sapi
C 5	410	41.2	5339	6	AX538620	AX538620 Sequence
C 6	410	41.2	5417	6	AX538619	AX538619 Sequence
C 7	410	41.2	5462	6	AX538621	AX538621 Sequence
C 8	410	41.2	8689	6	AX538622	AX538622 Sequence
C 9	410	41.2	11443	6	AX538624	AX538624 Sequence
C 10	410	41.2	12057	6	AX538627	AX538627 Sequence
C 11	410	41.2	12446	9	HSMDMR	X14298 Human mRNA
C 12	410	41.2	13957	6	AX409637	AX409637 Sequence
C 13	410	41.2	13957	6	AX538581	AX538581 Sequence
C 14	410	41.2	13957	9	HUMDYS	M18533 Homo sapien
C 15	396	39.8	2148	11	G15848	G15848 human STS C
C 16	396	39.8	2891	6	AX538618	AX538618 Sequence
C 17	359	36.0	13977	6	AR220819	AR220819 Sequence
C 18	350	35.1	350	9	AF213444	AF213444 Homo sapi
C 19	289	29.0	2110	9	HUMDMDX	M92650 Human Duche
C 20	278	27.9	2563	9	AK129855	AK129855 Homo sapi
C 21	260	26.1	3163	6	E30223	E30223 Shortened d
C 22	260	26.1	3172	6	E30222	E30222 Shortened d
C 23	260	26.1	4075	6	E30221	E30221 Shortened d
C 24	260	26.1	4402	6	E30219	E30219 Shortened d
C 25	260	26.1	4402	6	E30220	E30220 Shortened d
C 26	251	25.2	3747	6	E30218	E30218 Shortened d
C 27	231	23.2	212120	2	AC023414	AC023414 Homo sapi
C 28	218	21.9	218	9	HUMDYS20	M86903 H.sapiens d
C 29	146	14.7	2148	11	G15848	G15848 human STS C
C 30	146	14.7	2563	9	AK129855	AK129855 Homo sapi
C 31	146	14.7	2691	6	AX538618	AX538618 Sequence
C 32	146	14.7	4658	9	BC028720	BC028720 Homo sapi
C 33	146	14.7	8689	6	AX538622	AX538622 Sequence
C 34	146	14.7	11443	6	AX538624	AX538624 Sequence
C 35	146	14.7	13957	6	AX409637	AX409637 Sequence
C 36	146	14.7	13957	6	AX538581	AX538581 Sequence
C 37	146	14.7	13957	9	HUMDYS	M18533 Homo sapien
C 38	146	14.7	98056	9	AC006061	AC006061 Homo sapi
C 39	145	14.6	2005	6	AX817318	AX817318 Sequence
C 40	126	12.7	13977	6	AR220819	AR220819 Sequence
C 41	102	10.2	13887	4	AF070485	AF070485 Carls fam
C 42	94	9.4	189131	10	AL645477	AL645477 Mouse DNA
C 43	94	9.4	272578	2	AC108338	AC108338 Rattus no
C 44	94	9.4	279539	2	AC114184	AC114184 Rattus no
C 45	91	9.1	13815	6	AX306153	AX306153 Sequence

ALIGNMENTS

RESULT 1
AC006061/c
LOCUS
DEFINITION Homo sapiens X BAC GSHB-19024 (genome systems Human BAC Library)
complete sequence.
AC006061
AC006061.1 GI:4204246
VERSION HTG.
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 98056)
REFERENCE
AUTHORS Muzny,D., Arenson,A.D., Bouck,J., Brundage,E., Bunac,C., Chen,Z.,
Di,W., Ding,Y., Dugan,S., Durbin,J., Forcum,J., Garcia,C.,
98056 bp DNA linear PRI 01-MAY-2002

Correll, J.H., Correll, L.L., Hernandez, J., Jackson, L., Kondejewski, N., Leal, B., Lichtarge, O., Liu, W., Logan, O., Lu, J., Martinez, C., Oswal, G., Pampelli, L.R., Parish, B.J., Perez, L., Rashid, N.D., Rives, C., Scherer, S.E., Shen, H., Simon, M., Vo, Q., Williamson, A., Worley, K.C., Yu, W., Zhou, X., Nelson, D. and Gibbs, R.A.

Direct Submission
Unpublished
2 (bases 1 to 98056)
Worley, K.C.

Direct Submission
Submitted (26-NOV-1998) Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 98056)
Worley, K.C.

Direct Submission
Submitted (30-JAN-1999) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
4 (bases 1 to 98056)
Worley, K.C.

Direct Submission
Submitted (02-FEB-1999) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
5 (bases 1 to 98056)
Worley, K.C.

Direct Submission
Submitted (04-FEB-1999) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
6 (bases 1 to 98056)
Worley, K.C.

Direct Submission
Submitted (28-MAR-2001) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
7 (bases 1 to 98056)
Worley, K.C.

Direct Submission
Submitted (07-MAR-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
8 (bases 1 to 98056)
Worley, K.C.

Direct Submission
Submitted (01-MAY-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Jan 30, 1999 this sequence version replaced gi:4176317.
INFORMATION: <http://gc.bcm.tmc.edu:8088/home.html> or email gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:

STSs are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases.

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	/db_xref="taxon:9606"
	/chromosome="X"
	/clone="GSHB-19024"
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repeat_region	40..82
	/function="clone overlap"
repeat_region	378..507
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repeat_region	518..570
	/rpt_family="L1MD3"
repeat_region	571..729
	/complement(518..570)
	/rpt_family="(GGA)n"
repeat_region	730..987
	/rpt_family="L1MC4"
repeat_region	991..1400
	/rpt_family="MSTD"
repeat_region	1649..1779
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repeat_region	1783..2212
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repeat_region	4713..5074
	/rpt_family="(GA)n"
repeat_region	5702..7108
	/rpt_family="THE1B"
repeat_region	8670..8734
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repeat_region	9069..9144
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repeat_region	9254..9351
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repeat_region	9537..9688
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repeat_region	9797..9942
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repeat_region	10403..10429
	/rpt_family="WIR"
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repeat_region	12556..13643
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repeat_region	13618..13808
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repeat_region	14327..14625
	/complement(13997..14297)
repeat_region	14648..14915
	/rpt_family="MER4D"
repeat_region	14915..15147
	/rpt_family="AluSg"
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	/rpt_family="MER4D"
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JOURNAL

Submitted (14-FEB-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Apr 11, 2000 this sequence version replaced gi:6970579.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html

COMMENT

----- Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WtBR
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence_submissions@genome.wi.mit.edu
 ----- Project Information
 Center project name: L6731
 Center clone name: 767_B_5
 ----- Summary Statistics
 Sequencing vector: M13; M77815; 100% of reads
 Chemistry: Dye-terminator Big Dye; 100% of reads
 Assembly program: Phrap; version 0.960731
 Consensus quality: 185633 bases at least Q40
 Consensus quality: 195115 bases at least Q30
 Consensus quality: 201837 bases at least Q20
 Insert size: 178000; agarose-fp
 Insert size: 209120; sum-of-contigs
 Quality coverage: 5.8 in Q20 bases; agarose-fp
 Quality coverage: 4.9 in Q20 bases; sum-of-contigs

 * NOTE: This is a 'working draft' sequence. It currently consists of 31 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

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1144 2224: contig of 1081 bp in length
2225 2324: gap of 100 bp
2325 3345: contig of 1021 bp in length
3346 3445: gap of 100 bp
3446 4502: contig of 1057 bp in length
4503 4603: gap of 100 bp
4603 5758: contig of 1154 bp in length
5759 5858: gap of 100 bp
5859 7097: contig of 1241 bp in length
7098 7197: gap of 100 bp
7198 8236: contig of 1039 bp in length
8237 9353: gap of 100 bp
9354 9453: gap of 100 bp
9454 10705: contig of 1252 bp in length
10706 10805: gap of 100 bp
10806 12332: contig of 1527 bp in length
12333 12433: gap of 100 bp
12434 13877: contig of 1445 bp in length
13878 13977: gap of 100 bp
13978 15201: contig of 1224 bp in length
15202 15301: gap of 100 bp
15302 16968: contig of 1667 bp in length
16969 17069: gap of 100 bp
17070 19061: contig of 1992 bp in length
19062 19161: gap of 100 bp
19162 22639: contig of 3479 bp in length
22640 22740: gap of 100 bp
22741 23739: contig of 3637 bp in length
23740 24768: gap of 100 bp
24769 25275: contig of 2799 bp in length
25276 26375: gap of 100 bp
26376 29376: contig of 3890 bp in length
29377 30366: gap of 100 bp
30367 31113: contig of 3647 bp in length
31114 33930: contig of 2278 bp in length
33931 37113:

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* 39391 39490: gap of 100 bp
* 39491 43630: contig of 4200 bp in length
* 43631 43790: gap of 100 bp
* 43791 47621: contig of 3831 bp in length
* 47622 47721: gap of 100 bp
* 47722 53852: contig of 6131 bp in length
* 53853 53952: gap of 100 bp
* 53953 63432: contig of 9470 bp in length
* 63433 63522: gap of 100 bp
* 63523 77361: contig of 13839 bp in length
* 77362 77461: gap of 100 bp
* 77462 92465: contig of 15004 bp in length
* 92466 106951: contig of 14386 bp in length
* 106952 107051: gap of 100 bp
* 107052 129678: contig of 22627 bp in length
* 129679 151427: contig of 21649 bp in length
* 151428 151527: gap of 100 bp
* 151528 181838: contig of 30311 bp in length
* 181839 181939: gap of 100 bp
* 181940 212120: contig of 30182 bp in length.

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              /db_xref="taxon:9606"
              /chromosome="14"
              /map="14"
              /clone="RP11-767B5"
              /clone_lib="RPC1-11 Human Male BAC"
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              /note="assembly_fragment"
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          misc_feature      2325..3345
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Db	2189	AAATAAATCATATATTTTGTGAAGGGTAGTGGTATATATCTAGTATCTTACAGTAGTATCTTCT	2248
Qy	690	AAAGTCGTGTTATGTTTGTGTTAAACAATGGCAGGTTTTTACAGCTCTATGCAATTTGTACAAA	749
Db	2249	AAAGTCGTGTTATGTTTGTGTTAAACAATGGCAGGTTTTTACAGCTCTATGCAATTTGTACAAA	2308
Qy	750	AAAGTTATAAGAAAACACTACATGTAATAATCTTGATAGCTTAATAACTTGCCTATTTCTTTATA	809
Db	2309	AAAGTTATAAGAAAACACTACATGTAATAATCTTGATAGCTTAATAACTTGCCTATTTCTTTATA	2368
Qy	810	TGGAACGCATTTTGGGTGTTGTTTAAAAATTTTAAACAGTTTAAAGAAGA	859
Db	2369	TGGAACGCATTTTGGGTGTTGTTTAAAAATTTTAAACAGTTTAAAGAAGA	2418
RESULT 4			
LOCUS	BC028720	4658 bp mRNA linear	PRI 25-AUG-2003
DEFINITION	Homo sapiens dystrophin (muscular dystrophy, Duchenne and Becker types), transcript variant Dp71b, mRNA (CDNA clone IMAGE:4822807), complete cds.		
ACCESSION	BC028720		
VERSION	BC028720.1	GI:20379675	
KEYWORDS	Homo sapiens (human)		
SOURCE	Homo sapiens		
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
REFERENCE	1 (bases 1 to 4658)		
AUTHORS	Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G., Klausner, R.D., Collins, F.S., Wagner, L.H., Shenman, C.M., Schuler, G.D., Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K., Hopkins, R.F., Jordan, H., Moore, I., Max, S.I., Wang, J., Haieh, F., Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L., Stapleton, L., Soares, M.B., Bonaldo, M.F., Casavant, T.L., Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S., Carninci, P., Brown, C., Raha, S.S., Loquellano, N.A., Peters, G.J., Abramson, R.D., Mallek, S.J., Bosak, S.A., McEwan, P.J., McKernan, K.J., Mulek, J.A., Gunaratne, P.H., Richards, S., Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W., Villalón, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A., Fanny, J., Helton, E., Kettman, M., Madan, A., Rodrigues, S., Sanchez, A., Whitting, M., Madan, A., Young, A.C., Shevchenko, Y., Bouffard, G.G., Blakeley, R.W., Touchman, J.W., Green, E.D., Dickson, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myers, R.M., Butlerfield, Y.S., Krzywicki, M.I., Skalska, U., Smalls, D.E., Schnerch, A., Schein, J.E., Jones, S.J. and Marra, M.A.		
TITLE	Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences		
JOURNAL	Proc. Natl. Acad. Sci. U.S.A.	99 (26), 16899-16903	(2002)
MEDLINE	22388257		
PUBMED	12477932		
REFERENCE	2 (bases 1 to 4658)		
AUTHORS	Strausberg, R.		
TITLE	Direct Submission		
JOURNAL	Submitted (29-APR-2002) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590.		
REMARK	USA		
COMMENT	NIH-MGC Project URL: http://mgc.nci.nih.gov Contact: MGC help desk Email: cgapsb-remail.nih.gov Tissue Procurement: Miklos Palkovits, M.D., Ph.D. cDNA Library Preparation: Michael J. Brownstein (NHGRI) & Shiraki Toshiyuki and Piero Carninci (RIKEN) cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Institute for Systems Biology http://www.systemsbiology.org contact: anadan@systemsbiology.org Anup Madan, Jessica Fahay, Erin Helton, Mark Kettman, Anuradha Madan, Stephanie Rodriguez, Amy Sanchez and Michelle Whitting		

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Series: IRAK Plate: 46 Row: j Column: 3
This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 5032296
This clone has the following problem: The cds is short compared to the longest cds in the locus.

FEATURES
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/clone_lib="NIH_MGC_97"
/lab_host="DH10B"
/note="Vector: pBluescript"
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gene

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1. 458
/notes="DMD"
/gene/synonyms: BMD, DXS142, DXS164, DXS206, DXS230,
DXS239, DXS268, DXS269, DXS270, DXS272"
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/db xref="MIM:300377"
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CDS

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/db_xref="LocusID:1756"
/taxation="MREOULGHETOTCWDHPKQMTLSLADLNINVFSAYRTAMKL
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CDGRRLHLHDSIPIROLGVSAGSGNSLECEPIGRSLSEKHFNDICQSFPGG
LEFSVMVLPVLRHVAASATKHQAKCNICEPIGRSLSEKHFNDICQSFPGG
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misc: feature

041. 1562
/note=ZZ; Region: Zinc finger, ZZ type. Zinc finger
present in dystrophin, CBP/p300. ZZ in dystrophin binds
calmodulin Putative zinc finger"
Ab_mval="CDD:pfam00559"

ORIGIN

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Query Match 41.2%; Score 410; DB 9; Length 4658;
Best Local Similarity 100.0%; Pred. No. 5.3e+203;
Mismatched 41.0; Conservative 0; Mismatches 0;
Gaps 0;

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150 ACCACCAATCTACGAGTCTTTCCACATGGCAGATGATTGGGCAGAGCGGATGGAGTC 509

CCGACCTCAGTGGATGTTGGTCAGACCGATCGACTC
1028 AGGCACAACTAGGAAGACCTTTCATCATCGACATGATTGGCAGACCGATCGACTC
450

DB
193B AGGACACATGTAGGAAGTCTTTCCACATGCCAGATGATTTCGGCAGCGGCGTGCTGCTCTT

510 GGGGACTGTCACCTCATCAACAATGCAGATGAGAAGGAGGAGAATAAATGTTTTTACACTCTCTGATT 569

[illegible]

DB 1998 CTTAGTATCAGTCATGACAGAATGAAGAGGGAAACAGATTGTTCTTATGCTCTCCCGGTA 629

570 CCGCATGGTTTATTAATAAATTCATACACCAAGAGGATTAGACATTAAGAATTAACTG
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Db
2038 CCCGATGGTATTATAAATTCATACAGCAAAAGAGA^aTAGAACAGAAAAGAGTTCTCCTTG
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630 AAATAAATCTATAATTGTCGAGGGTAGTGGTAATATACGTGAAGAATTCAGATGAATTC
QY

Db
2118 AAATAAATCTATATTTTGTGAGGGGTAGTGGTATTAATACGAGAGAAATTCAGTAGATTCCT 2119

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Qy	810	TGGAACGCATTTTGGGTTGTTTAAAAATTTTAAACAGTTTATAAAGAAAGA	859
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RESULT 5			
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LOCUS	Sequence 40 from Patent WO0229056.		linear
DEFINITION	AX538620		
ACCESSION	AX538620.1	GI:25271166	
VERSION			
KEYWORDS	synthetic construct		
SOURCE	synthetic construct		
ORGANISM	artificial sequences.		
REFERENCE	1		
AUTHORS	Chamberlain, J.S. and Harper, S.O.		
TITLE	Mini-dysprophic nucleic acid and peptide sequences		
JOURNAL	Patent: WO 0229056-A 40 11-APR-2002;		
	THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)		
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Best Local Similarity	100.0%;	Pred No. 5.2e-203;	
Matches 410;	Conservative 0;	Mismatches 0;	Indels 0; Gaps 0;
Qy	450	AGGACACAATGTAGAAAGTCTTTTCCACATGGCAGATGATTGGGACAGCGATGGAGTC	509
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Qy	510	CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTACAACTCCTGATT	569
Db	4551	CTTAGTATCAGTCATGACAGATGAAGAGGAGCAGATAAATGTTTACAACTCCTGATT	4610
Qy	570	CCCGCATGGTTTTATATATTTTCATACACAAGAGGATTAGACAGTAAGAGTTTACAAAG	629
Db	4611	CCCGCATGGTTTTATATATTTTCATACACAAGAGGATTAGACAGTAAGAGTTTACAAAG	4670
Qy	630	AAATAAATCTATATTTTGTGAGGGTAGTGTTATATCTGTAGATTTCAGTAGTTTCT	689
Db	4671	AAATAAATCTATATTTTGTGAGGGTAGTGTTATATCTGTAGATTTCAGTAGTTTCT	4730
Qy	690	AAGTCTGTTATGTTTCTTAAATATGTCAGGTTTACACGTCCTATGCAATGTACAAAA	749
Db	4731	AAGTCTGTTATGTTTCTTAAATATGTCAGGTTTACACGTCCTATGCAATGTACAAAA	4799
Qy	750	AAGTTATAGAAAACTACATGTAAATCTTGATAGCTAAATAACTTGCCTTTCTTTATA	809
Db	4791	AAGTTATAGAAAACTACATGTAAATCTTGATAGCTAAATAACTTGCCTTTCTTTATA	4855
Qy	810	TGGACCGCATTTGGGTTGTTTAAAAATTTTAAACAGTTTATAAAGAAAGA	859
Db	4851	TGGACCGCATTTGGGTTGTTTAAAAATTTTAAACAGTTTATAAAGAAAGA	4900
RESULT 6			
AX538619	AX538619	5417 bp	DNA
LOCUS	Sequence 39 from Patent WO0229056.		linear
DEFINITION	AX538619		
ACCESSION	AX538619.1	GI:25271163	
VERSION			
KEYWORDS	synthetic construct		
SOURCE			

Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACTCCTGATT 569
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QY 570 CCCGATGGTTTTTATAATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
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QY 690 AAGTCGTGTTATTTGTTTAAACAATGGCAGGTTTTACACGTCATGCAATTTGTACAAA 749
 Db 4854 AAGTCGTGTTATTTGTTTAAACAATGGCAGGTTTTACACGTCATGCAATTTGTACAAA 4913

QY 750 AAGTTATAAGAAACTACATGTAATACTTGTAGCTAAATAACTTGCATTTCTTTATA 809
 Db 4914 AAGTTATAAGAAACTACATGTAATACTTGTAGCTAAATAACTTGCATTTCTTTATA 4973

QY 810 TGGAAACGCAATTTGGGTTGTTTAAATAATTTATAACAGTTATAAAGAAAGA 859
 Db 4974 TGGAAACGCAATTTGGGTTGTTTAAATAATTTATAACAGTTATAAAGAAAGA 5023

RESULT 8
 AX538622
 LOCUS AX538622 8689 bp DNA linear PAT 23-NOV-2002
 DEFINITION Sequence 42 from Patent WO0229056.
 ACCESSION AX538622
 VERSION AX538622.1 GI:25271171
 KEYWORDS synthetic construct
 SOURCE synthetic construct
 ORGANISM synthetic construct
 REFERENCE 1
 AUTHORS Chamberlain, J.S. and Harper, S.O.
 TITLE Mini-dystrophin nucleic acid and peptide sequences
 JOURNAL Patent: WO 0229056-A 42 11-APR-2002;
 THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)
 FEATURES
 Location/Qualifiers
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ORIGIN
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 Best Local Similarity 100.0%; Pred. No. 4.9e-203;
 Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACTCCTGATT 569
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QY 570 CCCGATGGTTTTTATAATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
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QY 750 AAGTTATAAGAAACTACATGTAATACTTGTAGCTAAATAACTTGCATTTCTTTATA 809
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QY 810 TGGAAACGCAATTTGGGTTGTTTAAATAATTTATAACAGTTATAAAGAAAGA 859
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RESULT 9
 AX538624
 LOCUS AX538624 11443 bp DNA linear PAT 23-NOV-2002
 DEFINITION Sequence 44 from Patent WO0229056.
 ACCESSION AX538624
 VERSION AX538624.1 GI:25271175
 KEYWORDS synthetic construct
 SOURCE synthetic construct
 ORGANISM synthetic construct
 REFERENCE 1
 AUTHORS Chamberlain, J.S. and Harper, S.O.
 TITLE Mini-dystrophin nucleic acid and peptide sequences
 JOURNAL Patent: WO 0229056-A 44 11-APR-2002;
 THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)
 FEATURES
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 source 1..11443
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ORIGIN
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 Best Local Similarity 100.0%; Pred. No. 4.7e-203;
 Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 450 AGGACACAATGAGGAAGTCCTTTTCCACATGCGAGATGATTTGGCGAGCGATGGAGTC 509
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QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACTCCTGATT 569
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QY 810 TGGAAACGCAATTTGGGTTGTTTAAATAATTTATAACAGTTATAAAGAAAGA 859
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RESULT 10
 AX538627
 LOCUS AX538627 12057 bp DNA linear PAT 23-NOV-2002

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DEFINITION Sequence 47 from Patent WO0229056.
ACCESSION AX538627
VERSION AX538627.1 GI:25271181
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1
AUTHORS Chamberlain, J.S. and Harper, S.O.
TITLE Mini-dystrophin nucleic acid and peptide sequences
JOURNAL Patent: WO 0229056-A 47 11-APR-2002;
THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)
FEATURES
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QY 450 AGGACACAAATGAGGAAGTCTTTTCCACATGCGAGATGATTTGGCGAGCGATCGAGTC 509
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QY 570 CCCGATCGTGGTTTTATAATATTTATACACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
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DB 11574 TGGACCGCATTTTGGTGTGTTTAAAAATTTATAACAGTTATAAGAAAGA 11623
RESULT 11
HSDMDR 12446 bp mRNA linear PRI 12-SEP-1993
LOCUS Human mRNA for dystrophin.
DEFINITION X14298
ACCESSION X14298.1 GI:30845
VERSION Dmd gene; Duchenne muscular dystrophy; dystrophin.
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
    Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 12446)
AUTHORS Rosenthal, A., Speer, A., Billwitz, H., Cross, G.S., Forrest, S.M. and
    Davies, K.E.
TITLE Two human cDNA molecules coding for the Duchenne muscular dystrophy
    (DMD) locus are highly homologous
JOURNAL Nucleic Acids Res. 17 (13), 5391 (1989)
MEDLINE 89345106
PUBMED 2668885
REFERENCE 2 (bases 1 to 12446)
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AUTHORS Rosenthal, A.
TITLE Direct submission
JOURNAL Submitted (09-FEB-1989) Rosenthal A., Akademie der Wissenschaften
    der DDR, Zentralinstitut fuer Molekularbiologie, Robert-Roessle
    Str. 10, 1115 Berlin Buch, DDR
COMMENT see also M18533 and M20250 for Dmd seqs.; discrepancies compared to
    M18533 cDNA were located at x14298 pos. 496, 1772, 1965, 2449,
    3687, 4299, 4504, 5075, 5332, 5630 and 7194.
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ORIGIN

Query Match 41.2%; Score 410; DB 9; Length 12446;
Best Local Similarity 100.0%; Pred. No. 4.7e-203; Indels 0; Gaps 0;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 450 AGGACACAATGTAGGAAGTCCTTTCCACATGCGAGATGATTGGGCGAGCGATGGAGTC 509
Db 11143 AGGACACAATGTAGGAAGTCCTTTCCACATGCGAGATGATTGGGCGAGCGATGGAGTC 11202
QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACTCCTGATT 569
Db 11203 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACTCCTGATT 11262
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Db 11263 CCCGATGCTTTTATAATATTATACACAAAGAGGATTAGACAGTAAGAGTTTACAAG 11322
QY 630 AAATAAATCTATATTTTGTGAAGGTAGTGGTATTATATCTGTAGATTTCAGTAGTTCT 689
Db 11323 AAATAAATCTATATTTTGTGAAGGTAGTGGTATTATATCTGTAGATTTCAGTAGTTCT 11382
QY 690 AAGTCGTATTGTTTGTAAACAATGCGAGCTTTTACACGCTCATGCAATGTCACAAA 749
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QY 750 AAGTTATAGAAAACACTACATGTAATAATCTTGATAGCTAAATAACTTGCATTCTTTATA 809
Db 11443 AAGTTATAGAAAACACTACATGTAATAATCTTGATAGCTAAATAACTTGCATTCTTTATA 11502
QY 810 TGGACGCAATTTGGTGTCTTTTAAAAAATTATAACAGTTATAAAGAAAAGA 859
Db 11503 TGGACGCAATTTGGTGTCTTTTAAAAAATTATAACAGTTATAAAGAAAAGA 11552

RESULT 12
AX409637 13957 bp DNA linear PAT 14-JUN-2002
LOCUS
DEFINITION Sequence 2284 from Patent WO0229103.
ACCESSION AX409637
VERSION AX409637.1 GI:21442342
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
Alvares, C., Horne, D., Peres-da-Silva, S. and Vockley, J. G.
Gene expression profiles in liver cancer
Patent: WO 0229103-A 2284 11-APR-2002;
GENE LOGIC INC (US)
FEATURES
Location/Qualifiers
source
1. .13957
/organism="Homo sapiens"
/mol_type="unassigned DNA"
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/note="EMBL/GenBank Accession No. M18533"

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Query Match 41.2%; Score 410; DB 6; Length 13957;
Best Local Similarity 100.0%; Pred. No. 4.6e-203; Indels 0; Gaps 0;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY 690 AAGTCGTATTGTTTGTAAACAATGCGAGCTTTTACACGCTCATGCAATGTCACAAA 749
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QY 750 AAGTTATAGAAAACACTACATGTAATAATCTTGATAGCTAAATAACTTGCATTCTTTATA 809
Db 11553 AAGTTATAGAAAACACTACATGTAATAATCTTGATAGCTAAATAACTTGCATTCTTTATA 11612
QY 810 TGGACGCAATTTGGTGTCTTTTAAAAAATTATAACAGTTATAAAGAAAAGA 859
Db 11613 TGGACGCAATTTGGTGTCTTTTAAAAAATTATAACAGTTATAAAGAAAAGA 11662

RESULT 13
AX538581 13957 bp DNA linear PAT 23-NOV-2002
LOCUS
DEFINITION Sequence 1 from Patent WO0229056.
ACCESSION AX538581
VERSION AX538581.1 GI:25271086
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
Chamberlain, J. S. and Harper, S. Q.
Mini-dystrophin nucleic acid and peptide sequences
Patent: WO 0229056-A 1 11-APR-2002;
JOURNAL THE REGENTS OF THE UNIVERSITY OF MICHIGAN (US)
FEATURES
Location/Qualifiers
source
1. .13957
/organism="Homo sapiens"
/mol_type="unassigned DNA"
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Query Match 41.2%; Score 410; DB 6; Length 13957;
Best Local Similarity 100.0%; Pred. No. 4.6e-203; Indels 0; Gaps 0;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 450 AGGACACAATGTAGGAAGTCCTTTCCACATGCGAGATGATTGGGCGAGCGATGGAGTC 509
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QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACCTCCTGATT 569
Db 11313 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACCTCCTGATT 11372
QY 570 CCCGATGCTTTTATAATATTATACACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
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Db      11433 AAAATAAATCTATATTTTGTGAAGGTAGTGGTATATATACGTGATGATTCAGTAGTTCT 11492
Qy      690 AAGTCTGTTATTTGTTTAAACATGCGAGGTTTACACGCTCATGCAATTGTACAAA 749
Db      11493 AAGTCTGTTATTTGTTTAAACATGCGAGGTTTACACGCTCATGCAATTGTACAAA 11552
Qy      750 AAGTATTAAGAAACTACATGTAATCTTATAGTCAATTAATGCTGCTGCTTTTATA 809
Db      11553 AAGTATTAAGAAACTACATGTAATCTTATAGTCAATTAATGCTGCTGCTTTTATA 11612
Qy      810 TCGAACGCAATTTGGTGTGTTTAAAAATTTATACAGTATAAAGAAAGA 859
Db      11613 TCGAACGCAATTTGGTGTGTTTAAAAATTTATACAGTATAAAGAAAGA 11662

RESULT 14
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LOCUS      HUMDYS      13957 bp      mRNA      linear      PRI 25-MAY-2000
DEFINITION Homo sapiens dystrophin (DMD) mRNA, complete cds.
ACCESSION M18533 M17154 M18026 M20250
VERSION    M18533.1 GI:181856
KEYWORDS
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 1599)
AUTHORS   Koenig,M., Hoffman,B.P., Bertelson,C.J., Monaco,A.P., Feener,C. and
            Kunkel,L.M.
TITLE     Complete cloning of the Duchenne muscular dystrophy (DMD) cDNA and
            preliminary genomic organization of the DMD gene in normal and
            affected individuals
JOURNAL   Cell 50 (3), 509-517 (1987)
MEDLINE   87273512
PubMed    3607877
REFERENCE 2 (bases 1678 to 3830)
AUTHORS   Hoffman,E.P., Monaco,A.P., Feener,C.C. and Kunkel,L.M.
TITLE     Conservation of the Duchenne muscular dystrophy gene in mice and
            humans
JOURNAL   Science 238 (4825), 347-350 (1987)
MEDLINE   88018015
PubMed    3659917
REFERENCE 3 (bases 1 to 13957)
AUTHORS   Koenig,M., Monaco,A.P. and Kunkel,L.M.
TITLE     The complete sequence of dystrophin predicts a rod-shaped
            cytoskeletal protein
JOURNAL   Cell 53 (2), 219-226 (1988)
MEDLINE   88194521
PubMed    3282674
COMMENT   On May 25, 2000 this sequence version replaced gi:340693.
            Draft entry and computer-readable sequence kindly provided by
            M.Koenig, 01-APR-1988 The severity of muscular dystrophy is
            determined by the size of the deleted DNA segment. Deletions found
            in different patients were from positions 302-2200, 473-1168,
            1691-1810, and 1169-3011.
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            /note="G00-119-850"
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QITSLAQGYERTSSPKPRFKSYATQAAVYVTSPTSPSPSOHLEAPDKSFGSSL
MSEVNLDRYQALBEVLSWLSAEDTLQAQGEISNDVEVVDQVSHTEGYNMIDLTAH
QSRVNLQGLGSLIGTKLSDEBETEVOEONNLNSWECURQVHTEGYNMIDLTAH
DLNQKLELNDLTKTEERTKMEEPGLDLEKQVQHQHVLQDLEQEVNVN
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Query Match 41.2%; Score 410; DB 9; Length 13957;
 Best Local Similarity 100.0%; Pred. No. 4.6e-203;
 Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 450 AGGACACATGATGAGGATCTTTTCCACATGCGAGATGATTTGGGCGAGCGCATGAGTC 509

Db 11253 AGGACACAATGTAGAAAGCTCTTTCCACATGCGCAGATGATTGGCAGAGCGATGGAGTC 11312

QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACTCCTGATT 569

Db 11313 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGATAAATGTTTACAACTCCTGATT 11372

QY 570 CCCGATCGTTTATATATATTTATACAAAGAGGATTAGACAGTAAGAGTTTACAAAG 629

Db 11373 CCCGATCGTTTATATATATTTATACAAAGAGGATTAGACAGTAAGAGTTTACAAAG 11432

QY 630 AAATAAATCTATATTTTGTGAAGGTTAGTGGTATTTATCTGTAGATTTCAAGTTTCT 689

Db 11433 AAATAAATCTATATTTTGTGAAGGTTAGTGGTATTTATCTGTAGATTTCAAGTTTCT 11492

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Db 11493 AGTCTGTTTATGTTTGTAAACATGCGCAGTTTACACCTCTATGCAATTTGACAAAA 11552

QY 750 AAGTTTATAGAAAACTACATGTAATCTTGATAGCTAAATAAAGTTCCTTTATA 809

Db 11553 AAGTTTATAGAAAACTACATGTAATCTTGATAGCTAAATAAAGTTCCTTTATA 11612

QY 810 TCGAACGCAATTTGGTGTGTTTAAAAATTTATAACAGTTATAAAGAAAGA 859

Db 11613 TCGAACGCAATTTGGTGTGTTTAAAAATTTATAACAGTTATAAAGAAAGA 11662

RESULT 15

G15848 2148 bp DNA linear STS 19-JAN-1996

LOCUS human STS CHLC.UTR_01924_M18533.P56108 clone UTR_01924_M18533,

DEFINITION sequence tagged site.

ACCESSION G15848

VERSION G15848.1 GI:1161737

KEYWORDS STS; STS sequence; primer; sequence tagged site.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo. 1 (bases 1 to 2148)

AUTHORS Murray J., Sheffield, V., Weber, J.L., Duyk, G. and Buetow, K.H.

TITLE Cooperative Human Linkage Center

JOURNAL Unpublished (1995)

COMMENT Synonyms: UTR_01924_M18533, CHLC.UTR_01924_M18533.T36152

Contact: Dr. Jeffrey C. Murray

UofI

The University of Iowa

Department of Pediatrics, Iowa City, IA 52242, USA

Tel: (319) 356-3508

Fax: (319) 356-3347

Email: jeff-murray@uiowa.edu

Primer A: AACGATTTGGTGTGTTTA

Primer B: GATATCACCCAAAAGGATG

STS size: 189

PCR Profile:

denature: 30 seconds at 94 degrees C

annealing: 75 seconds at 55 degrees C

extension: 15 seconds at 72 degrees C

PCR cycles: 27

extension: 6 minutes at 72 degrees C

Protocol:

Template: 30ng genomic DNA

Primer: each 1.5 pmole

dNTPs: each 200 uM

Taq Polymerase: 0.3 units

Total Vol: 10 ul

Buffer:

MgCl2: 1.5mM

KCl: 50mM

Tris: 10mM

pH: 8.3

Prepared with primer pairs derived from M18533.

Location/Qualifiers

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/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

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Best Local Similarity 100.0%; Pred. No. 1.3e-195;

Matches 396; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 GAAGTCTTTCCACATGCGCAGATGTTTGGCAGAGCGATGCGATCCTTAGTATCAGTCA 60

QY 524 TGACAGATGAAGAAGGAGCAGAAATAAATGTTTACAACTCCTGATTCGCGCATGTTTT 583

Db 61 TGACAGATGAAGAAGGAGCAGAAATAAATGTTTACAACTCCTGATTCGCGCATGTTTT 120

QY 584 ATATATTTATACAAAGAGGATTAGACAGTAAGAGTTTACAAAGAAATTAATCTATAT 643

Db 121 ATATATTTATACAAAGAGGATTAGACAGTAAGAGTTTACAAAGAAATTAATCTATAT 180

QY 644 TTTTGTGAAGGGTAGTGGTATTATCTGTAGATTTTCAGTAGTTTCTAAGTCTGTTATTGT 703

Db 181 TTTTGTGAAGGGTAGTGGTATTATCTGTAGATTTTCAGTAGTTTCTAAGTCTGTTATTGT 240

QY 704 TTTGTTAAACAATGCGCAGGTTTTACACGTCTATGCAATTTGTACAAAAAGTTATAGAAAA 763

Db 241 TTTGTTAAACAATGCGCAGGTTTTACACGTCTATGCAATTTGTACAAAAAGTTATAGAAAA 300

QY 764 CTACATGTAAATCTGTAGCTAAATAAATCTTGCCCATTTCTTTATATGGAACGCAATTTG 823

Db 301 CTACATGTAAATCTGTAGCTAAATAAATCTTGCCCATTTCTTTATATGGAACGCAATTTG 360

QY 824 GGTGTTTAAAAATTTATACAGTTTATAAAGAAAGA 859

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Search completed: April 6, 2004, 16:30:24

Job time : 4178.52 secs

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C 2	126	12.7	13977	4	US-09-484-970B-60 Sequence 60, Appl
C 3	91	9.1	19307	3	US-08-836-022A-10 Sequence 10, Appl
C 4	91	9.1	19307	3	US-09-427-048A-10 Sequence 10, Appl
5	42	4.2	19307	3	US-08-836-022A-10 Sequence 10, Appl
6	42	4.2	19307	3	US-09-427-048A-10 Sequence 10, Appl
7	23	2.3	434	3	US-08-943-731-127 Sequence 127, Appl
8	23	2.3	24183	3	US-08-943-731-3 Sequence 3, Appl
9	20	2.0	891	4	US-08-936-165A-90 Sequence 90, Appl
C 10	20	2.0	1201	3	US-09-374-64C-1 Sequence 1, Appl
C 11	19	1.9	4766	5	PCT-US93-07261-10 Sequence 10, Appl
C 12	19	1.9	9139	4	US-09-322-478-22 Sequence 22, Appl
C 13	18	1.8	25	2	US-08-676-782-4 Sequence 4, Appl
C 14	18	1.8	342	2	US-08-676-782-11 Sequence 11, Appl
C 15	18	1.8	393	1	US-08-248-505-1 Sequence 1, Appl
C 16	18	1.8	393	2	US-08-676-782-6 Sequence 6, Appl
17	18	1.8	750	4	US-09-107-532A-866 Sequence 866, Appl
18	18	1.8	889	2	US-08-935-886-7 Sequence 7, Appl
19	18	1.8	889	2	US-08-935-886-13 Sequence 13, Appl
20	18	1.8	1098	4	US-08-956-171B-23 Sequence 23, Appl
C 21	18	1.8	1349	2	US-08-676-782-10 Sequence 10, Appl
C 22	18	1.8	1824	4	US-09-489-039A-4969 Sequence 4969, Appl
23	18	1.8	1919	4	US-09-398-179-1 Sequence 1, Appl
C 24	18	1.8	1919	4	US-09-398-179-2 Sequence 2, Appl
C 25	18	1.8	1966	4	US-09-221-017B-964 Sequence 964, Appl
26	18	1.8	2076	4	US-09-134-001C-1838 Sequence 1838, Appl
C 27	18	1.8	2382	4	US-09-107-532A-2569 Sequence 2569, Appl


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QY 750 AAGTTATAGAAAACCTACATGTAAATCTGTAGTAACTAAATAAATGCGCATTTCTTTATA 809
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QY 810 TCGAACGCAATTTGGTGTGTTTAAATTTATAACAGTTATAAAGAAAGA 859
DB 11614 TCGAACGCAATTTGGTGTGTTTAAATTTATAACAGTTATAAAGAAAGA 11663

RESULT 2
US-09-484-970B-60/c
; Sequence 60, Application US/09484970B
; Patent No. 6426186
; GENERAL INFORMATION:
; APPLICANT: Jones, Karen A.
; APPLICANT: Volkmut, Wayne
; APPLICANT: Walker, Michael G.
; TITLE OF INVENTION: BONE REMODELING GENES
; FILE REFERENCE: PB-0014 US
; CURRENT APPLICATION NUMBER: US/09/484,970B
; CURRENT FILING DATE: 2000-01-18
; NUMBER OF SEQ ID NOS: 172
; SOFTWARE: PERL Program
; SEQ ID NO 60
; LENGTH: 13977
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc.feature
; OTHER INFORMATION: Incyte ID No. 6426186 229357.11CB1
; NAME/KEY: unsure
; LOCATION: 11721-11761, 12294, 13969
; OTHER INFORMATION: a, t, c, g, or other
US-09-484-970B-60

Query Match 12.7%; Score 126; DB 4; Length 13977;
Best Local Similarity 100.0%; Pred. No. 2.5e-53;
Matches 126; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 851 AAAGAAGAAATTTATAAGGAAAGAAAGAAATACGCAATGCGCAAGTGGTGAAGCTGTGAA 910
DB 13315 AAAGAAGAAATTTATAAGGAAAGAAAGAAATACGCAATGCGCAAGTGGTGAAGCTGTGAA 13256
QY 911 CTCAGTGTGCAATATTCAGGAACACCCCAACCAAGTGAAGTGAATACATGA 970
DB 13255 CTCAGTGTGCAATATTCAGGAACACCCCAACCAAGTGAAGTGAATACATGA 13196
QY 971 GAAGCC 976
DB 13195 GAAGCC 13190

RESULT 3
US-08-836-022A-10/c
; Sequence 10, Application US/08836022A
; Patent No. 6001557
; GENERAL INFORMATION:
; APPLICANT: Trustees of the University of Pennsylvania
; APPLICANT: Wilson, James M.
; APPLICANT: Fisher, Krishna J.
; APPLICANT: Chen, Shu-Jen
; APPLICANT: Weitzman, Matthew
; TITLE OF INVENTION: Improved Adenovirus Virus and
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Howson and Howson
; STREET: Spring House Corporate Cntr, P O Box 457
; CITY: Spring House
; STATE: Pennsylvania
; COUNTRY: USA
; ZIP: 19477
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/427,048A
; FILING DATE: 21-Oct-1999
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; PRIOR APPLICATION NUMBER: 08/836,022
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Bak, Mary E.
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
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COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/836,022A
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA: US 08/331,381
; APPLICATION NUMBER: 28-OCT-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Bak, Mary E.
; REGISTRATION NUMBER: 31,215
; REFERENCE/DOCKET NUMBER: GNVEN.008PCT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 215-540-9200
; TELEFAX: 215-540-5818
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 19307 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: unknown
; MOLECULE TYPE: cdna
US-08-836-022A-10

Query Match 9.1%; Score 91; DB 3; Length 19307;
Best Local Similarity 100.0%; Pred. No. 8.4e-36;
Matches 91; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 665 TATACGTAGATTCAGTAGCTTCTTAAGTCTGTATTTGTTTAAATGCGCAGTTT 724
DB 3225 TATACGTAGATTCAGTAGCTTCTTAAGTCTGTATTTGTTTAAATGCGCAGTTT 3166
QY 725 TACAGTCTATGCAATTGTACAAAAAAGTTA 755
DB 3165 TACAGTCTATGCAATTGTACAAAAAAGTTA 3135

RESULT 4
US-09-427-048A-10/c
; Sequence 10, Application US/09427048A
; Patent No. 6203975
; GENERAL INFORMATION:
; APPLICANT: Trustees of the University of Pennsylvania
; APPLICANT: Wilson, James M.
; APPLICANT: Fisher, Krishna J.
; APPLICANT: Chen, Shu-Jen
; APPLICANT: Weitzman, Matthew
; TITLE OF INVENTION: Improved Adenovirus Virus and
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Howson and Howson
; STREET: Spring House Corporate Cntr, P O Box 457
; CITY: Spring House
; STATE: Pennsylvania
; COUNTRY: USA
; ZIP: 19477
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/427,048A
; FILING DATE: 21-Oct-1999
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; PRIOR APPLICATION NUMBER: 08/836,022
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Bak, Mary E.
```

```

US-08-836-022A-10
Query Match      4.2%; Score 42; DB 3; Length 19307;
Best Local Similarity 100.0%; Pred. No. 2.8e-11;
Matches 42; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      934 AACACCCCAAAACCAAGTGAGGTAGAGTAAGTACGATGAGAAGC 975
      |||
Db      1563 AACACCCCAAAACCAAGTGAGGTAGAGTAAGTACGATGAGAAGC 1604
      |||

RESULT 6
US-09-427-048A-10
; Sequence 10, Application US/09427048A
; Patent No. 6203575
; GENERAL INFORMATION:
; APPLICANT: Trustees of the University of Pennsylvania
; Wilson, James M.
; Fisher, Krishna J.
; Chen, Shu-Jen
; Weitzman, Matthew
; TITLE OF INVENTION: Improved Adenovirus Virus and
; Methods of Use Thereof
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Howson and Howson
; STREET: Spring House Corporate Cntr, P O Box 457
; CITY: Spring House
; STATE: Pennsylvania
; COUNTRY: USA
; ZIP: 19477
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/427,048A
; FILING DATE: 21-Oct-1999
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/836,022
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Bak, Mary E.
; REGISTRATION NUMBER: 31,215
; REFERENCE/DOCKET NUMBER: GNVPN.008PCT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 215-540-9200
; TELEFAX: 215-540-5818
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 19307 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: unknown
; MOLECULE TYPE: cDNA
; SEQUENCE DESCRIPTION: SEQ ID NO: 10:
; US-09-427-048A-10

Query Match      4.2%; Score 42; DB 3; Length 19307;
Best Local Similarity 100.0%; Pred. No. 2.8e-11;
Matches 42; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      934 AACACCCCAAAACCAAGTGAGGTAGAGTAAGTACGATGAGAAGC 975
      |||
Db      1563 AACACCCCAAAACCAAGTGAGGTAGAGTAAGTACGATGAGAAGC 1604
      |||

RESULT 7
US-08-943-731-127
; Sequence 127, Application US/08943731
; Patent No. 6265157

```


TITLE OF INVENTION: Polypeptides and Their Uses

NUMBER OF SEQUENCES: 534

CORRESPONDENCE ADDRESS:

ADDRESSEE: SmithKline Beecham Corporation

STREET: 709 Swedeland Road

CITY: King of Prussia

STATE: PA

COUNTRY: USA

ZIP: 19406-0939

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FASEQ for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/936,165A

FILING DATE: 24-SEP-1997

CLASSIFICATION: 536

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 60/027,032

FILING DATE: 24-SEP-1996

ATTORNEY/AGENT INFORMATION:

NAME: Gimmil, Edward R

REGISTRATION NUMBER: 38,891

REFERENCE/DOCKET NUMBER: P50549

TELECOMMUNICATION INFORMATION:

TELEPHONE: 610-270-4478

TELEFAX: 610-270-5090

TELEX:

INFORMATION FOR SEQ ID NO: 90:

SEQUENCE CHARACTERISTICS:

LENGTH: 891 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: Genomic DNA

US-08-936-165A-90

Query Match 2.0%; Score 20; DB 4; Length 891;

Best Local Similarity 100.0%; Pred. No. 2.7;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 686 TTCTAAGTCGTGTTATGTTT 705

Db 579 TTCTAAGTCGTGTTATGTTT 598

RESULT 10

US-09-274-642-1/c

Sequence 1, Application US/09274642A

Patent No. 6071729

GENERAL INFORMATION:

APPLICANT: Jeffries, Thomas W.

APPLICANT: Shi, Nian-Qing

TITLE OF INVENTION: Disruption of cytochrome c gene in xylose-fermenting yeast

FILE REFERENCE: cytochrome c

CURRENT APPLICATION NUMBER: US/09/274,642A

CURRENT FILING DATE: 1999-03-23

EARLIER APPLICATION NUMBER: 60/080,493

EARLIER FILING DATE: 1998-04-02

NUMBER OF SEQ ID NOS: 42

SOFTWARE: PatentIn Ver. 2.0

SEQ ID NO 1

LENGTH: 1201

TYPE: DNA

ORGANISM: Fichia stipitis

FEATURE:

NAME/KEY: CDS

LOCATION: (610)...(942)

US-09-274-642-1

Query Match

Best Local Similarity 2.0%; Score 20; DB 3; Length 1201;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 623 TTACAAGAAATAAATCTATA 642

Db 1062 TTACAAGAAATAAATCTATA 1043

RESULT 11

PCT-US93-07261-10

Sequence 10, Application PC/TUS9307261

GENERAL INFORMATION:

TITLE OF INVENTION: PfEMP3 MALARIA ANTIGEN, ANALOGS, ANTIBODIES AND USES THEREOF

NUMBER OF SEQUENCES: 23

CORRESPONDENCE ADDRESS:

ADDRESSEE: John H. C. Blasdale

STREET: One Giralda Farms

CITY: Madison

STATE: New Jersey

COUNTRY: USA

ZIP: 07940-1000

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: Apple Macintosh

OPERATING SYSTEM: Macintosh 6.0.5

SOFTWARE: Microsoft Word 5.1a

CURRENT APPLICATION DATA:

APPLICATION NUMBER: PCT/US93/07261

FILING DATE: 19930805

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07/927,531

FILING DATE: 07-AUG-1992

ATTORNEY/AGENT INFORMATION:

NAME: Blasdale, John H. C.

REGISTRATION NUMBER: 31,895

REFERENCE/DOCKET NUMBER: DX0288K

TELECOMMUNICATION INFORMATION:

TELEPHONE: 201-822-7398

TELEFAX: 201-822-7039

INFORMATION FOR SEQ ID NO: 10:

SEQUENCE CHARACTERISTICS:

LENGTH: 4766 base pairs

TYPE: nucleic acid

STRANDEDNESS: double

TOPOLOGY: linear

MOLECULE TYPE: cDNA to mRNA

ORIGINAL SOURCE:

ORGANISM: Plasmodium falciparum

STRAIN: Malayan Camp

IMMEDIATE SOURCE:

CLONE: p2b1p12-1

FEATURE:

NAME/KEY: CDS

LOCATION: 3..4766

PCT-US93-07261-10

Query Match 1.9%; Score 19; DB 5; Length 4766;

Best Local Similarity 100.0%; Pred. No. 8.8;

Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 864 TAAAGGAAAAAGAAATAA 882

Db 190 TAAAGGAAAAAGAAATAA 208

RESULT 12

US-09-322-478-22/c

Sequence 22, Application US/09322478

Patent No. 6331662

GENERAL INFORMATION:

APPLICANT: Wright, David A.

APPLICANT: Voytas, Daniel F.

TITLE OF INVENTION: Plant Retroelements and Methods Related Thereto

FILE REFERENCE: P-1065 ISURF Plant Retroelement

;; CURRENT APPLICATION NUMBER: US/09/322,478
;; CURRENT FILING DATE: 1999-05-28
;; EARLIER APPLICATION NUMBER: 60/087125
;; EARLIER FILING DATE: 1998-05-29
;; NUMBER OF SEQ ID NOS: 41
;; SOFTWARE: Patent in Ver. 2.0
;; SEQ ID NO 22
;; LENGTH: 9139
;; TYPE: DNA
;; ORGANISM: Glycine max
US-09-322-478-22

Query Match 1.8%; Score 19; DB 4; Length 9139;
Best Local Similarity 100.0%; Pred. No. 8.9;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 528 AGATGAAGAAGGAGCAGAA 546
DB 5801 AGATGAAGAAGGAGCAGAA 5783

RESULT 13
US-08-676-782-4/c
; Sequence 4, Application US/08676782
; Patent No. 5976792
; GENERAL INFORMATION:
; APPLICANT: CHEUNG, Ambrose
; APPLICANT: FISCHETTI, Vincent A.
; TITLE OF INVENTION: REGULATION OF EXOPROTEIN IN
; TITLE OF INVENTION: STAPHYLOCOCCUS AUREUS
; NUMBER OF SEQUENCES: 18
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BURNS, DOANE, SWECKER & MATHIS, L.L.P.
; STREET: P.O. Box 1404
; CITY: Alexandria
; STATE: Virginia
; COUNTRY: United States
; ZIP: 22313-1404
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/676,782
; FILING DATE: 08-JUL-1996
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/248,505
; FILING DATE: 25-MAY-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: McGowan, Malcolm K.
; REGISTRATION NUMBER: 39,300
; REFERENCE/DOCKET NUMBER: 016921-092
; TELEPHONE: (703) 836-6620
; TELEFAX: (703) 836-2021
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 25 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-08-676-782-4

Query Match 1.8%; Score 18; DB 2; Length 25;
Best Local Similarity 100.0%; Pred. No. 25;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 431 TTCTTTCTCTTTGTTTC 448
DB 24 TTCTTTCTCTTTGTTTC 7

RESULT 14
US-08-676-782-11/c
; Sequence 11, Application US/08676782
; Patent No. 5976792
; GENERAL INFORMATION:
; APPLICANT: CHEUNG, Ambrose
; APPLICANT: FISCHETTI, Vincent A.
; TITLE OF INVENTION: REGULATION OF EXOPROTEIN IN
; TITLE OF INVENTION: STAPHYLOCOCCUS AUREUS
; NUMBER OF SEQUENCES: 18
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BURNS, DOANE, SWECKER & MATHIS, L.L.P.
; STREET: P.O. Box 1404
; CITY: Alexandria
; STATE: Virginia
; COUNTRY: United States
; ZIP: 22313-1404
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/676,782
; FILING DATE: 08-JUL-1996
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/248,505
; FILING DATE: 25-MAY-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: McGowan, Malcolm K.
; REGISTRATION NUMBER: 39,300
; REFERENCE/DOCKET NUMBER: 016921-092
; TELEPHONE: (703) 836-6620
; TELEFAX: (703) 836-2021
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 342 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 157
; OTHER INFORMATION: /note= "Nucleotide at position 157
; OTHER INFORMATION: is N wherein N = C or T."
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 340
; OTHER INFORMATION: /note= "Nucleotide at position 340
; OTHER INFORMATION: is N wherein N = C or T."
US-08-676-782-11

Query Match 1.8%; Score 18; DB 2; Length 342;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 431 TTCTTTCTCTTTGTTTC 448
DB 150 TTCTTTCTCTTTGTTTC 133

RESULT 15
US-08-548-505-1/c
; Sequence 1, Application US/08248505
; Patent No. 5587288
; GENERAL INFORMATION:
; APPLICANT: CHEUNG, Ambrose
; APPLICANT: FISCHETTI, Vincent A.

TITLE OF INVENTION: REGULATION OF EXOPROTEIN IN
 TITLE OF INVENTION: STAPHYLOCOCCUS AUREUS
 NUMBER OF SEQUENCES: 5
 CORRESPONDENCE ADDRESS:
 ADDRESSES: Burns, Doane, Swecker & Mathis
 STREET: P.O. Box 1404
 CITY: Alexandria
 STATE: Virginia
 COUNTRY: United States
 ZIP: 22313-1404
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC Compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/248,505
 FILING DATE: 24-MAY-1994
 CLASSIFICATION: 435
 ATTORNEY/AGENT INFORMATION:
 NAME: Crane-Feury, Sharon E
 REGISTRATION NUMBER: 36,113
 REFERENCE/DOCKET NUMBER: 016921-018
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (703) 836-6620
 TELEFAX: (703) 836-2021
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 393 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)
 FEATURE:
 NAME/KEY: CDS
 LOCATION: 19..390
 US-08-248-505-1

Query Match 1.8%; Score 18; DB 1; Length 393;
 Best Local Similarity 100.0%; Pred. No. 27;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 431 TTCCTTCTCTTTGTTTC 448
 Db 168 TTCCTTCTCTTTGTTTC 151

Search completed: April 6, 2004, 17:42:28
 Job time: 111.385 secs


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QY 61 CAATCACTCATAGCCAGGTGGAAGAAGATGATCCCATCATGGAATATTCCTGTTCTGAT 120
Db 61 CAATCACTCATAGCCAGGTGGAAGAAGATGATCCCATCATGGAATATTCCTGTTCTGAT 120
QY 121 AGAATCTTGTGCTTATCTATGGAATCTTTTGTATATATTTACATGGGAACCTGAAT 180
Db 121 AGAATCTTGTGCTTATCTATGGAATCTTTTGTATATATTTACATGGGAACCTGAAT 180
QY 181 GTAGCTTGACATTTTCCATGTAAACACACAGTAGCTGATCCAACTAAAGCTGATCTA 240
Db 181 GTAGCTTGACATTTTCCATGTAAACACACAGTAGCTGATCCAACTAAAGCTGATCTA 240
QY 241 ACAACAACAGTGTATGCTTCAATTAAGAGCTTGTCTTCTGCGAAGCTGGTGA 300
Db 241 ACAACAACAGTGTATGCTTCAATTAAGAGCTTGTCTTCTGCGAAGCTGGTGA 300
QY 301 AATCAAACTTGTGCTGATACACCTCGATGAGCTTCTGCTGTCTTCAACCCAGAAATG 360
Db 301 AATCAAACTTGTGCTGATACACCTCGATGAGCTTCTGCTGTCTTCAACCCAGAAATG 360
QY 361 GGAATGATTTCCCAATGGGCAAGAAACAGAGTATGCTATCTATCTGCACTTTTGT 420
Db 361 GGAATGATTTCCCAATGGGCAAGAAACAGAGTATGCTATCTATCTGCACTTTTGT 420
QY 421 AAGTCTGCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCT 480
Db 421 AAGTCTGCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCT 480
QY 481 GCAGATGATTTGGGAGAGCGATGAGTCCCTTAGTATCAGTCATGACAGATGAAGAGGA 540
Db 481 GCAGATGATTTGGGAGAGCGATGAGTCCCTTAGTATCAGTCATGACAGATGAAGAGGA 540
QY 541 GCAGATGATTTGGGAGAGCGATGAGTCCCTTAGTATCAGTCATGACAGATGAAGAGGA 600
Db 541 GCAGATGATTTGGGAGAGCGATGAGTCCCTTAGTATCAGTCATGACAGATGAAGAGGA 600
QY 601 AAGAGGATTTAGACAGTGAAGGTTTCAAGAAATAAATCTATATTTTGTGAAGGTAGT 660
Db 601 AAGAGGATTTAGACAGTGAAGGTTTCAAGAAATAAATCTATATTTTGTGAAGGTAGT 660
QY 661 GTATTATAGTGTAGATTTTCAAGTGTCTTCAAGTGTCTTCAAGTGTCTTCAAGTGTCT 720
Db 661 GTATTATAGTGTAGATTTTCAAGTGTCTTCAAGTGTCTTCAAGTGTCTTCAAGTGTCT 720
QY 721 GTTTTACAGCTCTATGCAATTTTACAAAAGTTTATAAGAAACTACATGTAATCTTG 780
Db 721 GTTTTACAGCTCTATGCAATTTTACAAAAGTTTATAAGAAACTACATGTAATCTTG 780
QY 781 ATAGCTAAATTAATCTGCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCT 840
Db 781 ATAGCTAAATTAATCTGCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCT 840
QY 841 TAACAGTTTAAAGAAAGATTTATAAGGAAAGAAAGAAATTAACGCAATGGCAAGTGTG 900
Db 841 TAACAGTTTAAAGAAAGATTTATAAGGAAAGAAAGAAATTAACGCAATGGCAAGTGTG 900
QY 901 AGCTGTGAACCTGAGTGTGCAATTTATCAGGAACACCCCAAAACCAAGTGAAGTGA 960
Db 901 AGCTGTGAACCTGAGTGTGCAATTTATCAGGAACACCCCAAAACCAAGTGAAGTGA 960
QY 961 AATAGCATGAGAGCCGCTGTTGATGTTAAAT 996
Db 961 AATAGCATGAGAGCCGCTGTTGATGTTAAAT 996

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RESULT 2
 US-10-149-736-40
 ; Sequence 40, Application US/10149736
 ; Publication No. US20030216332A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Chamberlain, Jeffrey S.
 ; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences

```

; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10149,736
; CURRENT FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: Patent in version 3.1
; SEQ ID NO 40
; LENGTH: 5339
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
; US-10-149-736-40

Query Match 41.2%; Score 410; DB 15; Length 5339;
Best Local Similarity 100.0%; Pred. No. 2.6e-189;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 450 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGGAGAGCGATGAGTC 509
Db 4491 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGGAGAGCGATGAGTC 4550
QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGAAATAAATGTTTACAACTCCTGATT 569
Db 4551 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGAAATAAATGTTTACAACTCCTGATT 4610
QY 570 CCGCATGGTTTTTAAATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
Db 4611 CCGCATGGTTTTTAAATATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG 4670
QY 630 AATAAATCTATATTTTGTGAAGGCTAGTGTATTATATCTAGATTTCAGTAGTTTCT 689
Db 4671 AATAAATCTATATTTTGTGAAGGCTAGTGTATTATATCTAGATTTCAGTAGTTTCT 4730
QY 690 AAGTCTGTATTGTTTGTAAACAATGGCAGGTTTACAGTCTATGCAATTTGACAAA 749
Db 4731 AAGTCTGTATTGTTTGTAAACAATGGCAGGTTTACAGTCTATGCAATTTGACAAA 4790
QY 750 AGTTTATAGAAACTACATGTAATAATCTTGTAGCTAATAAATGTCCTATTTCTTTATA 809
Db 4791 AGTTTATAGAAACTACATGTAATAATCTTGTAGCTAATAAATGTCCTATTTCTTTATA 4850
QY 810 TGAAGCGCAATTTGGGTTGTTTAAATAATTTATAACAGTTTATAAGAAAGA 859
Db 4851 TGAAGCGCAATTTGGGTTGTTTAAATAATTTATAACAGTTTATAAGAAAGA 4900

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RESULT 3
 US-10-149-736-39
 ; Sequence 39, Application US/10149736
 ; Publication No. US20030216332A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Chamberlain, Jeffrey S.
 ; APPLICANT: Harper, Scott Q.
 ; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
 ; FILE REFERENCE: UM-06968
 ; CURRENT APPLICATION NUMBER: US/10149,736
 ; CURRENT FILING DATE: 2002-06-17
 ; PRIOR APPLICATION NUMBER: PCT/US01/31126
 ; PRIOR FILING DATE: 2001-10-04
 ; PRIOR APPLICATION NUMBER: 60/238,848
 ; PRIOR FILING DATE: 2000-10-06
 ; NUMBER OF SEQ ID NOS: 96
 ; SOFTWARE: Patent in version 3.1
 ; SEQ ID NO 39
 ; LENGTH: 5417
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Synthetic

US-10-149-736-39

Query Match 41.2%; Score 410; DB 15; Length 5417;
Best Local Similarity 100.0%; Pred. No. 2.6e-189; Mismatches 0; Indels 0; Gaps 0;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 450 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 509
DB 4569 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 4628

QY 510 CTTAGTATCAGTCATGACAGATGAGAGGAGCAGATTAATGTTTACAACTCTTGATT 569
DB 4629 CTTAGTATCAGTCATGACAGATGAGAGGAGCAGATTAATGTTTACAACTCTTGATT 4688

QY 570 CCCGATGGTGTATTTATATATTTCTATACAAAGAGGATTTAGACAGTAAGAGTTTACAAG 629
DB 4689 CCCGATGGTGTATTTATATATTTCTATACAAAGAGGATTTAGACAGTAAGAGTTTACAAG 4748

QY 630 AAATAAATCTATATTTTGTGAAGGTAGTGGTATTTATCTGTAGATTTCAAGTATTTCT 689
DB 4749 AAATAAATCTATATTTTGTGAAGGTAGTGGTATTTATCTGTAGATTTCAAGTATTTCT 4808

QY 690 AAGTCTGTATTTTGTAAACATGGCAGGTTTACACGCTCTATGCAATTTGACAAA 749
DB 4809 AAGTCTGTATTTTGTAAACATGGCAGGTTTACACGCTCTATGCAATTTGACAAA 4868

QY 750 AAGTTAAGAAAACACTACATGTAATCTTGATAGCTTAATAACTTGCATTTCTTTATA 809
DB 4869 AAGTTAAGAAAACACTACATGTAATCTTGATAGCTTAATAACTTGCATTTCTTTATA 4928

QY 810 TGGACGCAATTTGGTGTGTTTAAATAATTTATAACAGTTATAAGAAAGA 859
DB 4929 TGGACGCAATTTGGTGTGTTTAAATAATTTATAACAGTTATAAGAAAGA 4978

RESULT 4

US-10-149-736-41
; Sequence 41, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; CURRENT FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 41
; LENGTH: 5462
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-10-149-736-41

Query Match 41.2%; Score 410; DB 15; Length 5462;
Best Local Similarity 100.0%; Pred. No. 2.6e-189; Mismatches 0; Indels 0; Gaps 0;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 450 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 509
DB 4614 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 4673

QY 510 CTTAGTATCAGTCATGACAGATGAGAGGAGCAGAAATAATGTTTACAACTCTTGATT 569
DB 4674 CTTAGTATCAGTCATGACAGATGAGAGGAGCAGAAATAATGTTTACAACTCTTGATT 4733

QY 570 CCCGATGGTGTATTTATATATTTCTATACAAAGAGGATTTAGACAGTAAGAGTTTACAAG 629

DB 4734 CCCGATGGTGTATTTATATATTTCTATACAAAGAGGATTTACACAGTAAGAGTTTACAAG 4793

QY 630 AAATAAATCTATATTTTGTGAAGGTAGTGGTATTTATCTGTAGATTTCAAGTATTTCT 689

DB 4794 AAATAAATCTATATTTTGTGAAGGTAGTGGTATTTATCTGTAGATTTCAAGTATTTCT 4853

QY 690 AAGTCTGTATTTTGTAAACATGGCAGGTTTACACGCTCTATGCAATTTGTAACAAA 749

DB 4854 AAGTCTGTATTTTGTAAACATGGCAGGTTTACACGCTCTATGCAATTTGTAACAAA 4913

QY 750 AAGTTAAGAAAACACTACATGTAATCTTGATAGCTTAATAACTTGCATTTCTTTATA 809

DB 4914 AAGTTAAGAAAACACTACATGTAATCTTGATAGCTTAATAACTTGCATTTCTTTATA 4973

QY 810 TGGACGCAATTTGGTGTGTTTAAATAATTTATAACAGTTATAAGAAAGA 859

DB 4974 TGGACGCAATTTGGTGTGTTTAAATAATTTATAACAGTTATAAGAAAGA 5023

RESULT 5

US-10-149-736-42
; Sequence 42, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; CURRENT FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 42
; LENGTH: 8689
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-10-149-736-42

Query Match 41.2%; Score 410; DB 15; Length 8689;
Best Local Similarity 100.0%; Pred. No. 2.6e-189; Mismatches 0; Indels 0; Gaps 0;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 450 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 509
DB 5985 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 6044

QY 510 CTTAGTATCAGTCATGACAGATGAGAGGAGCAGATTAATGTTTACAACTCTTGATT 569
DB 6045 CTTAGTATCAGTCATGACAGATGAGAGGAGCAGATTAATGTTTACAACTCTTGATT 6104

QY 570 CCCGATGGTGTATTTATATATTTCTATACAAAGAGGATTTAGACAGTAAGAGTTTACAAG 629
DB 6105 CCCGATGGTGTATTTATATATTTCTATACAAAGAGGATTTAGACAGTAAGAGTTTACAAG 6164

QY 630 AAATAAATCTATATTTTGTGAAGGTAGTGGTATTTATCTGTAGATTTCAAGTATTTCT 689

DB 6165 AAATAAATCTATATTTTGTGAAGGTAGTGGTATTTATCTGTAGATTTCAAGTATTTCT 6224

QY 690 AAGTCTGTATTTTGTAAACATGGCAGGTTTACACGCTCTATGCAATTTGTAACAAA 749

DB 6225 AAGTCTGTATTTTGTAAACATGGCAGGTTTACACGCTCTATGCAATTTGTAACAAA 6284

QY 750 AAGTTAAGAAAACACTACATGTAATCTTGATAGCTTAATAACTTGCATTTCTTTATA 809

DB 6285 AAGTTAAGAAAACACTACATGTAATCTTGATAGCTTAATAACTTGCATTTCTTTATA 6344

QY 810 TGGACGCAATTTGGTGTGTTTAAATAATTTAAACAGTTATAAAGAAAGA 859
DB 6345 TGGACGCAATTTGGTGTGTTTAAATAATTTAAACAGTTATAAAGAAAGA 6394

RESULT 6

US-10-149-736-44
; Sequence 44, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; APPLICANT: Harper, Scott Q.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; CURRENT FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 44
; LENGTH: 11443
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-10-149-736-44

Query Match 41.2%; Score 410; DB 15; Length 11443;
Best Local Similarity 100.0%; Pred. No. 2.7e-189;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 450 AGGACACAATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 509
DB 8739 AGGACACAATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 8798
QY 510 CTTAGTATCAGTCATGACAGATGAAGAGGACAGCAATAAATGTTTACAACCTCTGATT 569
DB 8799 CTTAGTATCAGTCATGACAGATGAAGAGGACAGCAATAAATGTTTACAACCTCTGATT 8858
QY 570 CCGCATGCTTATTTTGTGAAGGCTAGTGTATTACTGTAGATTTCAGTAGTTTCT 629
DB 8859 CCGCATGCTTATTTTGTGAAGGCTAGTGTATTACTGTAGATTTCAGTAGTTTCT 8918
QY 630 AATAAATCTATATTTTGTGAAGGCTAGTGTATTACTGTAGATTTCAGTAGTTTCT 689
DB 8919 AATAAATCTATATTTTGTGAAGGCTAGTGTATTACTGTAGATTTCAGTAGTTTCT 8978
QY 690 AAGTCTGTTATTGTTTGTGAAGGCTAGTGTATTACTGTAGATTTCAGTAGTTTCT 749
DB 8979 AAGTCTGTTATTGTTTGTGAAGGCTAGTGTATTACTGTAGATTTCAGTAGTTTCT 9038
QY 750 AGTTATAGAAACCTACATGTAATAATCTGTAGCTAATAACTTGCATTTCTTTATA 809
DB 9039 AGTTATAGAAACCTACATGTAATAATCTGTAGCTAATAACTTGCATTTCTTTATA 9098
QY 810 TGGACGCAATTTGGTGTGTTTAAATAATTTAAACAGTTATAAAGAAAGA 859
DB 9099 TGGACGCAATTTGGTGTGTTTAAATAATTTAAACAGTTATAAAGAAAGA 9148

RESULT 7

US-10-149-736-47
; Sequence 47, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; APPLICANT: Harper, Scott Q.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736

; CURRENT FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,848
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 47
; LENGTH: 12057
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-10-149-736-47

Query Match 41.2%; Score 410; DB 15; Length 12057;
Best Local Similarity 100.0%; Pred. No. 2.7e-189;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 450 AGGACACAATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 509
DB 11214 AGGACACAATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTC 11273
QY 510 CTTAGTATCAGTCATGACAGATGAAGAGGACAGCAATAAATGTTTACAACCTCTGATT 569
DB 11274 CTTAGTATCAGTCATGACAGATGAAGAGGACAGCAATAAATGTTTACAACCTCTGATT 11333
QY 570 CCGCATGCTTATTTTGTGAAGGCTAGTGTATTACTGTAGATTTCAGTAGTTTCT 629
DB 11334 CCGCATGCTTATTTTGTGAAGGCTAGTGTATTACTGTAGATTTCAGTAGTTTCT 11393
QY 630 AATAAATCTATATTTTGTGAAGGCTAGTGTATTACTGTAGATTTCAGTAGTTTCT 689
DB 11394 AATAAATCTATATTTTGTGAAGGCTAGTGTATTACTGTAGATTTCAGTAGTTTCT 11453
QY 690 AAGTCTGTTATTGTTTGTGAAGGCTAGTGTATTACTGTAGATTTCAGTAGTTTCT 749
DB 11454 AAGTCTGTTATTGTTTGTGAAGGCTAGTGTATTACTGTAGATTTCAGTAGTTTCT 11513
QY 750 AAGTTATAGAAACCTACATGTAATAATCTGTAGCTAATAACTTGCATTTCTTTATA 809
DB 11514 AAGTTATAGAAACCTACATGTAATAATCTGTAGCTAATAACTTGCATTTCTTTATA 11573
QY 810 TGGACGCAATTTGGTGTGTTTAAATAATTTAAACAGTTATAAAGAAAGA 859
DB 11574 TGGACGCAATTTGGTGTGTTTAAATAATTTAAACAGTTATAAAGAAAGA 11623

RESULT 8

US-09-782-378A-22
; Sequence 22, Application US/09782378A
; Patent No. US20020102731A1
; GENERAL INFORMATION:
; APPLICANT: Hearing, Patrick
; APPLICANT: Bahou, Wadie
; APPLICANT: Sandalon, Ziv
; APPLICANT: Gratenko, Dmitri
; TITLE OF INVENTION: Adenoviral Vectors
; FILE REFERENCE: STONYB-04970
; CURRENT APPLICATION NUMBER: US/09/782,378A
; CURRENT FILING DATE: 2001-02-12
; PRIOR APPLICATION NUMBER: 60/237,747
; PRIOR FILING DATE: 2000-10-02
; NUMBER OF SEQ ID NOS: 27
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 22
; LENGTH: 13957
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-782-378A-22

Query Match 41.2%; Score 410; DB 9; Length 13957;
Best Local Similarity 100.0%; Pred. No. 2.7e-189;

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Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 450 AGGACACAAATGAGGAAGTCTTTTCCACATGCGAGATGATTGGCGAGCGATGGAGTC 509
DB 11253 AGGACACAAATGAGGAAGTCTTTTCCACATGCGAGATGATTGGCGAGCGATGGAGTC 11312
QY 510 CTTAGTATCAGTCATGACAGATGAAGAGGAGCGAGATGAATGTTTACAACTCCTGATT 569
DB 11313 CTTAGTATCAGTCATGACAGATGAAGAGGAGCGAGATGAATGTTTACAACTCCTGATT 11372
QY 570 CCCGATGCTGTTTATAATATTTATACACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
DB 11373 CCCGATGCTGTTTATAATATTTATACACAAAGAGGATTAGACAGTAAGAGTTTACAAG 11432
QY 630 AAATTAATCTATATTTTGTGAGGAGTGTGTTATATCTGATGTTTCAAGTCTTCT 689
DB 11433 AAATTAATCTATATTTTGTGAGGAGTGTGTTATATCTGATGTTTCAAGTCTTCT 11492
QY 690 AAGTCTGTTATTTGTTTAAACAATGCGCAGGTTTACACGCTCTATGCAATTTGTAACAAA 749
DB 11493 AAGTCTGTTATTTGTTTAAACAATGCGCAGGTTTACACGCTCTATGCAATTTGTAACAAA 11552
QY 750 AAGTTATAAGAAACTACATGTAATCTTGATAGCTAAATAAATCTTGCATTTCTTTATA 809
DB 11553 AAGTTATAAGAAACTACATGTAATCTTGATAGCTAAATAAATCTTGCATTTCTTTATA 11612
QY 810 TGGAACGCAATTTGGGTTGTTTAAATAATTTATAACAGTTTATAAGAAAGA 859
DB 11613 TGGAACGCAATTTGGGTTGTTTAAATAATTTATAACAGTTTATAAGAAAGA 11662
RESULT 9
US-09-880-107-2284
; Sequence 2284, Application US/09880107
; Patent No. US20020142981A1
; GENERAL INFORMATION:
; APPLICANT: Horne, Darci T.
; APPLICANT: Vockley, Joseph G.
; APPLICANT: Scherf, Uwe
; APPLICANT: Gene Logic, Inc.
; TITLE OF INVENTION: Gene Expression Profiles in Liver Cancer
; FILE REFERENCE: 44921-5028-WO
; CURRENT APPLICATION NUMBER: US/09/880,107
; CURRENT FILING DATE: 2001-06-14
; PRIOR APPLICATION NUMBER: US 60/211,379
; PRIOR FILING DATE: 2000-06-14
; PRIOR APPLICATION NUMBER: US 60/237,054
; PRIOR FILING DATE: 2000-10-02
; NUMBER OF SEQ ID NOS: 3950
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2284
; LENGTH: 13957
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: Genbank Accession No. US20020142981A1 M18533
US-09-880-107-2284
Query Match 41.2%; Score 410; DB 9; Length 13957;
Best Local Similarity 100.0%; Pred. No. 2.7e-189;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 450 AGGACACAAATGAGGAAGTCTTTTCCACATGCGAGATGATTGGCGAGCGATGGAGTC 509
DB 11253 AGGACACAAATGAGGAAGTCTTTTCCACATGCGAGATGATTGGCGAGCGATGGAGTC 11312
QY 510 CTTAGTATCAGTCATGACAGATGAAGAGGAGCGAGATGAATGTTTACAACTCCTGATT 569
DB 11313 CTTAGTATCAGTCATGACAGATGAAGAGGAGCGAGATGAATGTTTACAACTCCTGATT 11372
QY 570 CCCGATGCTGTTTATAATATTTATACACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
DB 11373 CCCGATGCTGTTTATAATATTTATACACAAAGAGGATTAGACAGTAAGAGTTTACAAG 11432
QY 630 AAATTAATCTATATTTTGTGAGGAGTGTGTTATATCTGATGTTTCAAGTCTTCT 689
DB 11433 AAATTAATCTATATTTTGTGAGGAGTGTGTTATATCTGATGTTTCAAGTCTTCT 11492
QY 690 AAGTCTGTTATTTGTTTAAACAATGCGCAGGTTTACACGCTCTATGCAATTTGTAACAAA 749
DB 11493 AAGTCTGTTATTTGTTTAAACAATGCGCAGGTTTACACGCTCTATGCAATTTGTAACAAA 11552
QY 750 AAGTTATAAGAAACTACATGTAATCTTGATAGCTAAATAAATCTTGCATTTCTTTATA 809
DB 11553 AAGTTATAAGAAACTACATGTAATCTTGATAGCTAAATAAATCTTGCATTTCTTTATA 11612
QY 810 TGGAACGCAATTTGGGTTGTTTAAATAATTTATAACAGTTTATAAGAAAGA 859
DB 11613 TGGAACGCAATTTGGGTTGTTTAAATAATTTATAACAGTTTATAAGAAAGA 11662
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QY 630 AAATTAATCTATATTTTGTGAGGAGTGTGTTATATCTGATGTTTCAAGTCTTCT 689
DB 11433 AAATTAATCTATATTTTGTGAGGAGTGTGTTATATCTGATGTTTCAAGTCTTCT 11492
QY 690 AAGTCTGTTATTTGTTTAAACAATGCGCAGGTTTACACGCTCTATGCAATTTGTAACAAA 749
DB 11493 AAGTCTGTTATTTGTTTAAACAATGCGCAGGTTTACACGCTCTATGCAATTTGTAACAAA 11552
QY 750 AAGTTATAAGAAACTACATGTAATCTTGATAGCTAAATAAATCTTGCATTTCTTTATA 809
DB 11553 AAGTTATAAGAAACTACATGTAATCTTGATAGCTAAATAAATCTTGCATTTCTTTATA 11612
QY 810 TGGAACGCAATTTGGGTTGTTTAAATAATTTATAACAGTTTATAAGAAAGA 859
DB 11613 TGGAACGCAATTTGGGTTGTTTAAATAATTTATAACAGTTTATAAGAAAGA 11662
RESULT 10
US-10-149-736-1
; Sequence 1, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; APPLICANT: Harper, Scott Q.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; CURRENT FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,948
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
; LENGTH: 13957
; TYPE: DNA
; ORGANISM: Homo sapiens
; OTHER INFORMATION:
US-10-149-736-1
Query Match 41.2%; Score 410; DB 15; Length 13957;
Best Local Similarity 100.0%; Pred. No. 2.7e-189;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 450 AGGACACAAATGAGGAAGTCTTTTCCACATGCGAGATGATTGGCGAGCGATGGAGTC 509
DB 11253 AGGACACAAATGAGGAAGTCTTTTCCACATGCGAGATGATTGGCGAGCGATGGAGTC 11312
QY 510 CTTAGTATCAGTCATGACAGATGAAGAGGAGCGAGATGAATGTTTACAACTCCTGATT 569
DB 11313 CTTAGTATCAGTCATGACAGATGAAGAGGAGCGAGATGAATGTTTACAACTCCTGATT 11372
QY 570 CCCGATGCTGTTTATAATATTTATACACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
DB 11373 CCCGATGCTGTTTATAATATTTATACACAAAGAGGATTAGACAGTAAGAGTTTACAAG 11432
QY 630 AAATTAATCTATATTTTGTGAGGAGTGTGTTATATCTGATGTTTCAAGTCTTCT 689
DB 11433 AAATTAATCTATATTTTGTGAGGAGTGTGTTATATCTGATGTTTCAAGTCTTCT 11492
QY 690 AAGTCTGTTATTTGTTTAAACAATGCGCAGGTTTACACGCTCTATGCAATTTGTAACAAA 749
DB 11493 AAGTCTGTTATTTGTTTAAACAATGCGCAGGTTTACACGCTCTATGCAATTTGTAACAAA 11552
QY 750 AAGTTATAAGAAACTACATGTAATCTTGATAGCTAAATAAATCTTGCATTTCTTTATA 809
DB 11553 AAGTTATAAGAAACTACATGTAATCTTGATAGCTAAATAAATCTTGCATTTCTTTATA 11612
QY 810 TGGAACGCAATTTGGGTTGTTTAAATAATTTATAACAGTTTATAAGAAAGA 859
DB 11613 TGGAACGCAATTTGGGTTGTTTAAATAATTTATAACAGTTTATAAGAAAGA 11662
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RESULT 11
US-10-342-887-434
; Sequence 434, Application US/10342887
; Publication No. US20040058340A1
; GENERAL INFORMATION:
; APPLICANT: Dai, Hongyue
; APPLICANT: He, Yudong
; APPLICANT: Linsley, Peter S.
; APPLICANT: Mao, Mao
; APPLICANT: Roberts, Christopher J.
; APPLICANT: Van 't Veer, Laura Johanna
; APPLICANT: Van de Vijver, Marc J.
; APPLICANT: Bernards, Rene
; TITLE OF INVENTION: Diagnosis and Prognosis of Breast Cancer Patients
; FILE REFERENCE: 9301-188-999
; CURRENT APPLICATION NUMBER: US/10/342,887
; PRIOR FILING DATE: 2003-01-15
; PRIOR APPLICATION NUMBER: 60/298,918
; PRIOR FILING DATE: 2001-06-18
; PRIOR APPLICATION NUMBER: 60/380,710
; PRIOR FILING DATE: 2002-05-14
; PRIOR APPLICATION NUMBER: 10/172,118
; PRIOR FILING DATE: 2002-06-14
; NUMBER OF SEQ ID NOS: 2699
; SEQ ID NO 434
; LENGTH: 14069
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-342-887-434

Query Match 41.2%; Score 410; DB 12; Length 14069;
Best Local Similarity 100.0%; Pred. No. 2.7e-189;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 450 AGGACACATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGCGAGCGATGGAGTC 509
DB 11365 AGGACACATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGCGAGCGATGGAGTC 11424
QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGAGGAGGAGTAATGTTTACAACCTCCTGATT 569
DB 11425 CTTAGTATCAGTCATGACAGATGAAGAAGAGGAGGAGTAATGTTTACAACCTCCTGATT 11484
QY 570 CCGCATGTTTATAAATTCATACAAAGAGGATTTAGACAGTAAAGCTTGTACAGTTTACAAG 629
DB 11485 CCGCATGTTTATAAATTCATACAAAGAGGATTTAGACAGTAAAGCTTGTACAGTTTACAAG 11544
QY 630 AATAAATCTATATTTTGTGAAGGCTAGTGGTATTATATCTGTAGATTTTCACTAGTTTCT 689
DB 11545 AATAAATCTATATTTTGTGAAGGCTAGTGGTATTATATCTGTAGATTTTCACTAGTTTCT 11604
QY 690 AGTCTGTTATTTTGTGTACAAATGGCAGGTTTACACGCTATGCAATTTGTACAAA 749
DB 11605 AGTCTGTTATTTTGTGTACAAATGGCAGGTTTACACGCTATGCAATTTGTACAAA 11664
QY 750 AAGTTATAAGAAACTACATGTAATAATCTTTGATAGCTAAATAAATTTGCCATTTCTTTATA 809
DB 11665 AAGTTATAAGAAACTACATGTAATAATCTTTGATAGCTAAATAAATTTGCCATTTCTTTATA 11724
QY 810 TGAAGCGCATTTGGGTTGTTTAAAAATTTATAACAGTTATAAGAAAGA 859
DB 11725 TGAAGCGCATTTGGGTTGTTTAAAAATTTATAACAGTTATAAGAAAGA 11774

RESULT 12
US-10-342-887-981
; Sequence 981, Application US/10342887
; Publication No. US20040058340A1
; GENERAL INFORMATION:
; APPLICANT: Dai, Hongyue
; APPLICANT: He, Yudong
; APPLICANT: Linsley, Peter S.
; APPLICANT: Mao, Mao
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; APPLICANT: Roberts, Christopher J.
; APPLICANT: Van 't Veer, Laura Johanna
; APPLICANT: Van de Vijver, Marc J.
; APPLICANT: Bernards, Rene
; TITLE OF INVENTION: Diagnosis and Prognosis of Breast Cancer Patients
; FILE REFERENCE: 9301-188-999
; CURRENT APPLICATION NUMBER: US/10/342,887
; CURRENT FILING DATE: 2003-01-15
; PRIOR APPLICATION NUMBER: 60/298,918
; PRIOR FILING DATE: 2001-06-18
; PRIOR APPLICATION NUMBER: 60/380,710
; PRIOR FILING DATE: 2002-05-14
; PRIOR APPLICATION NUMBER: 10/172,118
; PRIOR FILING DATE: 2002-06-14
; NUMBER OF SEQ ID NOS: 2699
; SEQ ID NO 981
; LENGTH: 14082
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-342-887-981

Query Match 41.2%; Score 410; DB 12; Length 14082;
Best Local Similarity 100.0%; Pred. No. 2.7e-189;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 450 AGGACACATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGCGAGCGATGGAGTC 509
DB 11378 AGGACACATGTAGGAAGTCTTTCCACATGGCAGATGATTTGGCGAGCGATGGAGTC 11437
QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGAGGAGGAGTAATAATGTTTACAACCTCCTGATT 569
DB 11438 CTTAGTATCAGTCATGACAGATGAAGAAGAGGAGGAGTAATAATGTTTACAACCTCCTGATT 11497
QY 570 CCGCATGTTTATAAATTCATACAAAGAGGATTTAGACAGTAAAGCTTGTACAGTTTACAAG 629
DB 11498 CCGCATGTTTATAAATTCATACAAAGAGGATTTAGACAGTAAAGCTTGTACAGTTTACAAG 11557
QY 630 AATAAATCTATATTTTGTGAAGGCTAGTGGTATTATATCTGTAGATTTTCACTAGTTTCT 689
DB 11558 AATAAATCTATATTTTGTGAAGGCTAGTGGTATTATATCTGTAGATTTTCACTAGTTTCT 11617
QY 690 AAGTCTGTTATTTGTTTGTAAATGGAATGGCAGGTTTACAGTCTATGCAATTTGTACAAA 749
DB 11618 AAGTCTGTTATTTGTTTGTAAATGGAATGGCAGGTTTACAGTCTATGCAATTTGTACAAA 11677
QY 750 AAGTTATAAGAAACTACATGTAATAATCTTTGATAGCTAAATAAATTTGCCATTTCTTTATA 809
DB 11678 AAGTTATAAGAAACTACATGTAATAATCTTTGATAGCTAAATAAATTTGCCATTTCTTTATA 11737
QY 810 TGAAGCGCATTTGGGTTGTTTAAAAATTTATAACAGTTATAAGAAAGA 859
DB 11738 TGAAGCGCATTTGGGTTGTTTAAAAATTTATAACAGTTATAAGAAAGA 11787

RESULT 13
US-10-341-434-108
; Sequence 108, Application US/10341434
; Publication No. US20030215835A1
; GENERAL INFORMATION:
; APPLICANT: Origene Technologies
; TITLE OF INVENTION: Differentially Regulated Prostate Cancer Genes
; FILE REFERENCE: 90 204 205 R1
; CURRENT APPLICATION NUMBER: US/10/341,434
; CURRENT FILING DATE: 2003-07-18
; PRIOR APPLICATION NUMBER: US 60/348,154
; PRIOR FILING DATE: 2002-01-15
; PRIOR APPLICATION NUMBER: US 60/348,119
; PRIOR FILING DATE: 2002-01-15
; NUMBER OF SEQ ID NOS: 238
; SOFTWARE: Patent in version 3.1
; SEQ ID NO 108
; LENGTH: 14082
; TYPE: DNA
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; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (703)..(11388)
; OTHER INFORMATION:
US-10-149-736-108

Query Match 41.2%; Score 410; DB 15; Length 14082;
Best Local Similarity 100.0%; Pred. No. 2.7e-189;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 450 AGGACACAATGAGGAGTCTTTCCACATGCGAGTATTTGGCAGAGCGATGGAGTC 509
DB 11378 AGGACACAATGAGGAGTCTTTCCACATGCGAGTATTTGGCAGAGCGATGGAGTC 11437
QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGAGGACGAATAAATGTTTACAACTCCTGATT 569
DB 11438 CTTAGTATCAGTCATGACAGATGAAGAAGAGGACGAATAAATGTTTACAACTCCTGATT 11497
QY 570 CCCGATGGTTTTATATATTTATACAAACAAGAGGATAGACAGTAAGATTACAG 629
DB 11498 CCCGATGGTTTTATATATTTATACAAACAAGAGGATAGACAGTAAGATTACAG 11557
QY 630 AAATAAATCTATATTTTGTGAAGGTAGTGTATTTACTGTAGATTTCAGTAGTTTCT 689
DB 11558 AAATAAATCTATATTTTGTGAAGGTAGTGTATTTACTGTAGATTTCAGTAGTTTCT 11617
QY 690 AAGTCGTATTTGTTTGTAAATGCGAGTTTACACGCTATGCAATGTACAAA 749
DB 11618 AAGTCGTATTTGTTTGTAAATGCGAGTTTACACGCTATGCAATGTACAAA 11677
QY 750 AAGTTATGAAGAACTACATGTAATCTTGATAGCTAAATACCTGCAATTCCTTATA 809
DB 11678 AAGTTATGAAGAACTACATGTAATCTTGATAGCTAAATACCTGCAATTCCTTATA 11737
QY 810 TGAACGCAATTTGGTTGTTTAAATAATTTAAACAGTTTAAAGAAAGA 859
DB 11738 TGAACGCAATTTGGTTGTTTAAATAATTTAAACAGTTTAAAGAAAGA 11787

RESULT 14
US-10-149-736-38
; Sequence 38, Application US/10149736
; Publication No. US20030216332A1
; GENERAL INFORMATION:
; APPLICANT: Chamberlain, Jeffrey S.
; TITLE OF INVENTION: Mini-Dystrophin Nucleic Acids and Peptide Sequences
; FILE REFERENCE: UM-06968
; CURRENT APPLICATION NUMBER: US/10/149,736
; PRIOR FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US01/31126
; PRIOR FILING DATE: 2001-10-04
; PRIOR APPLICATION NUMBER: 60/238,648
; PRIOR FILING DATE: 2000-10-06
; NUMBER OF SEQ ID NOS: 96
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 38
; LENGTH: 2691
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-149-736-38

Query Match 39.8%; Score 396; DB 15; Length 2691;
Best Local Similarity 100.0%; Pred. No. 1.7e-182;
Matches 396; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 464 GAAGCTTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTCTTAGTATCAGTCA 523
DB 1 GAAGCTTTTCCACATGGCAGATGATTTGGCAGAGCGATGGAGTCTTAGTATCAGTCA 60
QY 524 TGACAGATGAAGAAGAGAGAGCAATAAATGTTTAACTCCTGATTCCCGCATGCTTTT 583

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DB 61 TGACAGATGAAGAAGAGAGCAATAAATGTTTAACTCTCGATTCCGCGATGGTTTTT 120
QY 584 ATAATATTTCATACAAAGAGAGGATTAGACAGTAAGAGTTTACAAAGAAATAAATCTATAT 643
DB 121 ATAATATTTCATACAAAGAGAGGATTAGACAGTAAGAGTTTACAAAGAAATAAATCTATAT 180
QY 644 TTTTGTGAAGGTAGTGTATTTACTGTAGATTTCAGTAGTTTCTTAAGTCTGTTATTGT 703
DB 181 TTTTGTGAAGGTAGTGTATTTACTGTAGATTTCAGTAGTTTCTTAAGTCTGTTATTGT 240
QY 704 TTTGTTAAACAATGCGCAGGTTTACACGCTATGCAATTTGTCACAAAAAGCTTAAAGAAA 763
DB 241 TTTGTTAAACAATGCGCAGGTTTACACGCTATGCAATTTGTCACAAAAAGCTTAAAGAAA 300
QY 764 CTACATGTAATCTTGATGATGCTAAATAAATCTGCAATTTCTTTATATGGAAGCAATTTG 823
DB 301 CTACATGTAATCTTGATGATGCTAAATAAATCTGCAATTTCTTTATATGGAAGCAATTTG 360
QY 824 GGTGTTTAAATAATTTATAACAGTTTAAAGAAAGA 859
DB 361 GGTGTTTAAATAATTTATAACAGTTTAAAGAAAGA 396

RESULT 15
US-09-796-692-3505/c
; Sequence 3505, Application US/09796692
; Publication No. US20020198362A1
; GENERAL INFORMATION:
; APPLICANT: Gaiger, Alexander
; APPLICANT: Algate, Paul A.
; APPLICANT: Mannion, Jane
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE DETECTION, DIAGNOSIS AND THERAPY
; FILE REFERENCE: 2077, 001200
; CURRENT APPLICATION NUMBER: US/09/796,692
; CURRENT FILING DATE: 2001-03-01
; PRIOR APPLICATION NUMBER: 60/186,126
; PRIOR FILING DATE: 2000-03-01
; PRIOR APPLICATION NUMBER: 60/190,479
; PRIOR FILING DATE: 2000-03-17
; PRIOR APPLICATION NUMBER: 60/200,545
; PRIOR FILING DATE: 2000-04-27
; PRIOR APPLICATION NUMBER: 60/200,303
; PRIOR FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: 60/200,779
; PRIOR FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: 60/200,999
; PRIOR FILING DATE: 2000-05-01
; PRIOR APPLICATION NUMBER: 60/202,084
; PRIOR FILING DATE: 2000-05-04
; PRIOR APPLICATION NUMBER: 60/206,201
; PRIOR FILING DATE: 2000-05-22
; PRIOR APPLICATION NUMBER: 60/218,950
; PRIOR FILING DATE: 2000-07-14
; PRIOR APPLICATION NUMBER: 60/222,903
; PRIOR FILING DATE: 2000-08-03
; PRIOR APPLICATION NUMBER: 60/223,416
; PRIOR FILING DATE: 2000-08-04
; PRIOR APPLICATION NUMBER: 60/223,378
; PRIOR FILING DATE: 2000-08-07
; NUMBER OF SEQ ID NOS: 9597
; SOFTWARE: PastSeq for Windows Version 3.0
; SEQ ID NO 3505
; LENGTH: 430
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-796-692-3505

Query Match 14.7%; Score 146; DB 9; Length 430;
Best Local Similarity 100.0%; Pred. No. 1.3e-60;
Matches 146; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 851 AAAGAAAGATTATAAGGAAAAAGAAATAACGCAATGGACAAGTGGTGAAGCTGTGAA 910

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Db	296	AAAGGAAGATTATAAAGGAAAAGAAAATAACGCAATGGACAAGTGGTGAAGCTGTGAA	237
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Db	236	CTCAGGTGTGCACAAATTATCAGGRACACCCCAAAACCAAGTGAGGTAGTAATAGCATGA	177
Qy	971	GAAGCCGTGTTTGATGTTTAATTAAAT	996
Db	176	GAAGCCGTGTTTGATGTTTAATTAAAT	151

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Job time : 450.211 secs

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OM nucleic - nucleic search, using sw model

Run on: April 6, 2004, 11:42:46 ; Search time 472.067 Seconds
(without alignments)
8963.144 Million cell updates/sec

Title: US-09-966-264D-2
Perfect score: 996
Sequence: 1 gtgttgatgattagataaa.....gtgttgatgattagataatt 996

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 3373863 seqs, 2124099041 residues

Word size : 0

Total number of hits satisfying chosen parameters: 6747726

Minimum DB seq length: 0
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Post-processing: Listing first 45 summaries

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1: Geneseqn1980s.*
2: Geneseqn1990s.*
3: Geneseqn2000s.*
4: Geneseqn2001as.*
5: Geneseqn2001bs.*
6: Geneseqn2002s.*
7: Geneseqn2003as.*
8: Geneseqn2003bs.*
9: Geneseqn2003cs.*
10: Geneseqn2004s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	996	100.0	996	ABK86497	ABK86497 Human Apo
2	996	100.0	1230	ABK86462	ABK86462 Human Apo
3	996	100.0	1234	ABK86463	ABK86463 Human Apo
4	410	41.2	2463	ABT33375	ABT33375 NOVX DNA
5	410	41.2	5339	ABK81998	ABK81998 DNA encod
6	410	41.2	5417	ABK81997	ABK81997 DNA encod
7	410	41.2	5462	ABK81999	ABK81999 DNA encod
8	410	41.2	8689	ABK82000	ABK82000 DNA encod
9	410	41.2	11443	ABK82002	ABK82002 DNA encod
10	410	41.2	12923	1 AAN90338	AAN90338 Sequence
11	410	41.2	13957	6 ABK81959	ABK81959 cDNA enco
12	410	41.2	13957	6 ABT10904	ABT10904 Human bre
13	410	41.2	13957	6 ABN95786	ABN95786 Gene #228
14	410	41.2	13957	6 ABK81996	ABK81996 Human dys
15	396	39.8	2691	6 ABK81996	ABK81996 Human dys
16	259	36.0	13977	6 ABK81996	ABK81996 Human bon
17	260	26.1	3163	3 AA48571	AA48571 A rod sho
18	260	26.1	3172	3 AA48570	AA48570 A rod sho
19	260	26.1	4075	3 AA248569	AA248569 A rod sho
20	260	26.1	4402	3 AA248568	AA248568 A rod sho
21	260	26.1	4402	3 AA248567	AA248567 A rod sho
22	251	25.2	3747	3 AA248566	AA248566 A rod sho
23	158	15.9	158	6 ABK86469	ABK86469 Human apo

C 24	146	14.7	200	6	ABK86468	ABK86468 Human dys
C 25	146	14.7	455	9	ADD32504	ADD32504 Human mit
C 26	146	14.7	2691	6	ABK81996	ABK81996 Human dys
C 27	146	14.7	8689	6	ABK82000	ABK82000 DNA encod
C 28	146	14.7	11443	6	ABK82002	ABK82002 DNA encod
C 29	146	14.7	13957	6	ABK81959	ABK81959 cDNA enco
C 30	146	14.7	13957	6	ABT10904	ABT10904 Human bre
C 31	146	14.7	13957	6	ABN95786	ABN95786 Gene #228
C 32	146	14.7	13957	6	ABK81996	ABK81996 Human dys
C 33	145	14.6	2005	7	ABT33376	ABT33376 NOVX DNA
C 34	137	13.8	137	6	ABK86496	ABK86496 Human Apo
C 35	126	12.7	13977	6	ABK86496	ABK86496 Human bon
C 36	94	9.4	1044	6	ABK86464	ABK86464 Mouse gen
C 37	91	9.1	13815	2	AAV18885	AAV18885 Mus muscu
C 38	91	9.1	13815	6	ABK81960	ABK81960 cDNA enco
C 39	91	9.1	13815	6	ABT199799	ABT199799 Mouse isc
C 40	91	9.1	19307	2	AAT27558	AAT27558 Shuttle v
C 41	60	6.0	60	6	ABK86471	ABK86471 Human apo
C 42	46	4.6	959	6	ABN74601	ABN74601 Bovine em
C 43	42	4.2	42	6	ABK86474	ABK86474 Human apo
C 44	42	4.2	13815	2	AAV18885	AAV18885 Mus muscu
C 45	42	4.2	13815	6	ABK81960	ABK81960 cDNA enco

ALIGNMENTS

RESULT 1
ABK86497
ID ABK86497 standard; cDNA; 996 BP.
XX
AC ABK86497;
XX
DT 27-AUG-2002 (first entry)
XX
DE Human Apo-dystrophin-4 cDNA.
XX
KW Human; ss; gene: apo-dystrophin-4; inversion sequence; gene therapy;
KW protein truncation; muscular dystrophy; leukaemia.
XX
OS Homo sapiens.

Key Location/Qualifiers
CDS 1..996
/tag= a
/product= "Apo-dystrophin-4 protein appearing as AA098739"
FT FT /partial
FT FT /note= "No start or stop codon shown"
FT FT /transl_except= (pos:7..9,aa:Xaa)
FT FT /transl_except= (pos:49..51,aa:Xaa)
FT FT /transl_except= (pos:187..189,aa:Xaa)
FT FT /transl_except= (pos:202..204,aa:Xaa)
FT FT /transl_except= (pos:217..219,aa:Xaa)
FT FT /transl_except= (pos:253..255,aa:Xaa)
FT FT /transl_except= (pos:394..396,aa:Xaa)
FT FT /transl_except= (pos:547..549,aa:Xaa)
FT FT /transl_except= (pos:565..567,aa:Xaa)
FT FT /transl_except= (pos:616..618,aa:Xaa)
FT FT /transl_except= (pos:649..651,aa:Xaa)
FT FT /transl_except= (pos:682..684,aa:Xaa)
FT FT /transl_except= (pos:709..711,aa:Xaa)
FT FT /transl_except= (pos:790..792,aa:Xaa)
FT FT /transl_except= (pos:841..843,aa:Xaa)
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FT FT /transl_except= (pos:907..909,aa:Xaa)
FT FT /transl_except= (pos:952..954,aa:Xaa)
FT FT /transl_except= (pos:982..984,aa:Xaa)
FT FT /transl_except= (pos:988..990,aa:Xaa)
FT FT /note= "Xaa= unknown, encoded by in frame stop codon"

FT /transl_except= (pos:451..453,aa:Xaa)
 FT /transl_except= (pos:487..489,aa:Xaa)
 FT /transl_except= (pos:499..504,aa:Xaa-Xaa)
 FT /transl_except= (pos:628..630,aa:Xaa)
 FT /transl_except= (pos:781..783,aa:Xaa)
 FT /transl_except= (pos:799..801,aa:Xaa)
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 FT /transl_except= (pos:883..885,aa:Xaa)
 FT /transl_except= (pos:916..920,aa:Xaa)
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 FT /transl_except= (pos:1024..1026,aa:Xaa)
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 FT /transl_except= (pos:1084..1086,aa:Xaa)
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 FT /transl_except= (pos:1186..1188,aa:Xaa)
 FT /transl_except= (pos:1216..1218,aa:Xaa)
 FT /transl_except= (pos:1222..1224,aa:Xaa)
 FT /note= "Xaa= unknown, encoded by in frame stop codon"

GB2368064-A.

24-APR-2002.

16-JAN-2001; 2001GB-00001124.

30-SEP-2000; 2000US-0237079P.

(IMCR) IMPERIAL CANCER RES TECHNOLOGY LTD.

(BARB/) BARBER E.

Barber E;

WPI: 2002-429042/46.

P-PSDB; AAU98729.

New human regulatory polynucleotide, useful for treating disorders associated with protein truncation, particularly muscular dystrophy, and related peptides and antibodies.

Disclosure; Fig 4; 222pp; English.

The invention relates to a polynucleotide (I) comprising, or consisting of, apo-dystrophin-4 inversion sequence appearing as ABK86496, or its functional equivalents (e.g. the apo-dystrophin-4 cDNA sequence appearing as ABK86497). Also included are polynucleotides that hybridize to either strand of (I), a vector containing (I), a cell containing (I) or the vector, proteins and peptides encoded by (I), a protein homologous with human dystrophin that is expressed on cell surfaces in vivo antibodies (Ab) specific for the protein and method of screening for leukemia cells by analyzing DNA for presence of (I) or by detecting presence of (II). The apo-dystrophin-4 inversion sequence is a regulatory element that controls expression (transcription and translation) of associated DNA, and may allow read-through of stop codons. The apo-dystrophin-4 inversion sequence is used in gene therapy of diseases associated with truncation of proteins, particularly muscular dystrophy and also leukaemia, but more generally (I) is a regulatory sequence used to control expression of any attached gene. Analysis of DNA for (I), or detection of proteins (II) encoded by (I), can be used to screen for leukaemic cells and related diseases. Antibodies raised against (II) can be used therapeutically, to inhibit (II) activity, also to detect (II) in screening assays. The present sequence is the full length cDNA sequence for human apo-dystrophin-4 containing a plurality of stop codons some of which may be read through due to the presence of (I) in the apo-dystrophin-4 gene

Sequence 1230 BP; 404 A; 189 C; 259 G; 378 T; 0 U; 0 Other;

Query Match 100.0%; Score 996; DB 6; Length 1230;

Best Local Similarity 100.0%; Pred. No. 0;

Matches 996; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 GTGGTTTGATGTAGTAAAGAAATGTCGTTATACAGTAGAGAGTAATCAAT 60
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Db	235	GTGGTTTGATGTAGTAAAGAAATGTCGTTATACAGTAGAGAGTAATCAAT	294
Qy	61	CAATCACTCATGCCAAGGTGGAAGAGATGTATCCCATCATGGAATATTCCTGTTCTGAT	120
Db	295	CAATCACTCATGCCAAGGTGGAAGAGATGTATCCCATCATGGAATATTCCTGTTCTGAT	354
Qy	121	AGAAATCTTGCTTATCTATGGAATCTTTTGATATATATTTACATTTGGGACCTGAAT	180
Db	355	AGAAATCTTGCTTATCTATGGAATCTTTTGATATATATTTACATTTGGGACCTGAAT	414
Qy	181	GTAGCTTGACATTTTCCATGTAAACACCCAGTAGCTGATCCAAACATTAAGCTGATCTA	240
Db	415	GTAGCTTGACATTTTCCATGTAAACACCCAGTAGCTGATCCAAACATTAAGCTGATCTA	474
Qy	241	AAAAACAAGCTGTAATGGCTTCATTAAGGCTTTGCTTCTTCTGGAACCTGTTGAA	300
Db	475	AAAAACAAGCTGTAATGGCTTCATTAAGGCTTTGCTTCTTCTGGAACCTGTTGAA	534
Qy	301	AATCAAAACCTTGTGTGACACCTCGATCGAGCTCTCTGTGTGCTTCCACAGAAATG	360
Db	535	AATCAAAACCTTGTGTGACACCTCGATCGAGCTCTCTGTGTGCTTCCACAGAAATG	594
Qy	361	GGGAATGATTTCCAAATGGCAAGAAACAGAGTGATGCTATCTATCTGACCTTTTGA	420
Db	595	GGGAATGATTTCCAAATGGCAAGAAACAGAGTGATGCTATCTATCTGACCTTTTGA	654
Qy	421	AGTCTGTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCT	480
Db	655	AGTCTGTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCT	714
Qy	481	GCAGATGATTTGGCGAGCGAGTGAGTCTTATAGTATCAGTATGATGACAGATGAAGGA	540
Db	715	GCAGATGATTTGGCGAGCGAGTGAGTCTTATAGTATCAGTATGATGACAGATGAAGGA	774
Qy	541	GCAGATGATTTGGCGAGCGAGTGAGTCTTATAGTATCAGTATGATGACAGATGAAGGA	600
Db	775	GCAGATGATTTGGCGAGCGAGTGAGTCTTATAGTATCAGTATGATGACAGATGAAGGA	834
Qy	601	AAGAGGATTAAGACAGTAAGAGTTTACAAGAAATTAATCTATATTTTGTGAAGGTAGT	660
Db	835	AAGAGGATTAAGACAGTAAGAGTTTACAAGAAATTAATCTATATTTTGTGAAGGTAGT	894
Qy	661	GTATATATCTAGATTTTCAAGTATGATGATGATGATGATGATGATGATGATGATGATG	720
Db	895	GTATATATCTAGATTTTCAAGTATGATGATGATGATGATGATGATGATGATGATGATG	954
Qy	721	GTATATATCTAGATTTTCAAGTATGATGATGATGATGATGATGATGATGATGATGATG	780
Db	955	GTATATATCTAGATTTTCAAGTATGATGATGATGATGATGATGATGATGATGATGATG	1014
Qy	781	ATAGCTAAATTAATCTGCAATTTTATATGGAACGCAATTTTGGGTTGTTTAAAAATTA	840
Db	1015	ATAGCTAAATTAATCTGCAATTTTATATGGAACGCAATTTTGGGTTGTTTAAAAATTA	1074
Qy	841	TACAGTTTAAAGAAAGTAATTAAGGAAAGAAAGTAATTAAGGAAAGTAATTAAGGAA	900
Db	1075	TACAGTTTAAAGAAAGTAATTAAGGAAAGTAATTAAGGAAAGTAATTAAGGAAAGTA	1134
Qy	901	AAGCTGTGAATCTAGGTTGTCACAATTTATCAGGAACACCCCAAAACCAAAAGTGA	960
Db	1135	AAGCTGTGAATCTAGGTTGTCACAATTTATCAGGAACACCCCAAAACCAAAAGTGA	1194
Qy	961	AATGAGATGAGAGCCGTTGATGATGATGATGATGATGATGATGATGATGATGATGATG	996
Db	1195	AATGAGATGAGAGCCGTTGATGATGATGATGATGATGATGATGATGATGATGATGATG	1230

RESULT 3

ABK86463

ID ABK86463 standard; cDNA; 1234 BP.

XX AC

XX ABK86463;

DT	27-AUG-2002	(first entry)	
XX	Human Apo-dystrophin-4 full cDNA fragment #1.		
DE	Human; ss; apo-dystrophin-4; inversion sequence; gene therapy;		
XX	protein truncation; muscular dystrophy; leukaemia.		
KW			
KW			
OS	Homo sapiens.		
OS			
PN	GB2368064-A.		
XX			
PD	24-APR-2002.		
PF	16-JAN-2001; 2001GB-00001124.		
PF			
PR	30-SEP-2000; 2000US-0237079P.		
XX	(IMCR) IMPERIAL CANCER RES TECHNOLOGY LTD.		
PA	(BARB/) BARBER E.		
PI	Barber E;		
XX			
DR	WPI; 2002-429042/46.		
XX			
PT	New human regulatory polynucleotide, useful for treating disorders		
PT	associated with protein truncation, particularly muscular dystrophy, and		
PT	related peptides and antibodies.		
XX			
PS	Disclosure, Fig 9; 222pp; English.		
XX			
CC	The invention relates to a polynucleotide (I) comprising, or consisting		
CC	of, apo-dystrophin-4 inversion sequence appearing as ABK86496, or its		
CC	functional equivalents (e.g. the apo-dystrophin-4 cDNA sequence appearing		
CC	as ABK86497). Also included are polynucleotides that hybridise to either		
CC	strand of (I), a vector containing (I), a cell containing (I) or the		
CC	vector, proteins and peptides encoded by (I), a protein homologous with		
CC	human dystrophin that is expressed on cell surfaces in vivo antibodies		
CC	(Ab) specific for the protein and method of screening for leukemia cells		
CC	by analysing DNA for presence of (I) or by detecting presence of (II).		
CC	The apo-dystrophin-4 inversion sequence is a regulatory element that		
CC	controls expression (transcription and translation) of associated DNA,		
CC	and may allow read-through of stop codons. The apo-dystrophin-4 inversion		
CC	sequence is used in gene therapy of diseases associated with truncation		
CC	of proteins, particularly muscular dystrophy and also leukaemia, but more		
CC	generally (I) is a regulatory sequence used to control expression of any		
CC	attached gene. Analysis of DNA for (I), or detection of proteins (II)		
CC	encoded by (I), can be used to screen for leukemic cells and related		
CC	diseases. Antibodies raised against (II) can be used therapeutically, to		
CC	inhibit (II) activity, also to detect (II) in screening assays. The		
CC	present sequence is a human apo-dystrophin-4 fragment matching a region		
CC	of mouse genomic DNA in the Apo-dystrophin-4 gene region		
XX			
SQ	Sequence 1234 BP; 405 A; 190 C; 260 G; 379 T; 0 U; 0 Other;		
	Query Match 100.0%; Score 996; DB 6; Length 1234;		
	Best Local Similarity 100.0%; Pred. No. 0;		
	Matches 996; Conservative 0; Mismatches 0; Indels 0; Gaps 0		
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Db	239 GTGGTTTGATTGATAGTAAAAAATGTTCTGTTAATCAAGTAGAGTAAGTAATCAAT 298		
QY	61 CAATCACTCATAGCCAAAGGTGGAAGAAGATGTATCCCATCATGGAATATTCCTGTTCTGAT 120		
Db	299 CAATCACTCATAGCCAAAGGTGGAAGAAGATGTATCCCATCATGGAATATTCCTGTTCTGAT 358		
QY	121 AGAAATCTTGCTTATCTATCGAATTCCTTTTGATATATATTTACATTTGGGAACCTGGAAT 180		
Db	359 AGAAATCTTGCTTATCTATCGAATTCCTTTTGATATATTTACATTTGGGAACCTGGAAT 418		
QY	181 GTAGCTTGACATTTTTCATGTAAACACACAGTAGCCTGATCCCAACATTAAGCTGATACTA 240		
Db	419 GTAGCTTGACATTTTTCATGTAAACACACAGTAGCCTGATCCCAACATTAAGCTGATACTA 478		

parasitic infection; Alzheimer's disease; stroke; forensic biology; immunogen; non-human transgenic animal; gene therapy; gene; ds. Unidentified.

WO200281517-A2.

17-OCT-2002.

22-JAN-2002; 2002WO-US02064.

19-JAN-2001; 2001US-0262892P.

23-JAN-2001; 2001US-0263598P.

24-JAN-2001; 2001US-0263799P.

25-JAN-2001; 2001US-0264117P.

26-JAN-2001; 2001US-0264439P.

26-JAN-2001; 2001US-0264478P.

30-JAN-2001; 2001US-0263351P.

02-MAR-2001; 2001US-0272870P.

14-MAR-2001; 2001US-0275927P.

14-MAR-2001; 2001US-0275990P.

15-MAR-2001; 2001US-0276449P.

20-MAR-2001; 2001US-0277358P.

23-MAR-2001; 2001US-0278151P.

29-MAR-2001; 2001US-0279857P.

20-APR-2001; 2001US-0285140P.

20-APR-2001; 2001US-0285141P.

30-APR-2001; 2001US-0287484P.

17-MAY-2001; 2001US-0291701P.

08-JUN-2001; 2001US-0296960P.

10-JUL-2001; 2001US-0304353P.

12-JUL-2001; 2001US-0304886P.

09-AUG-2001; 2001US-0311289P.

13-AUG-2001; 2001US-0311975P.

16-AUG-2001; 2001US-0312937P.

18-OCT-2001; 2001US-0330227P.

29-NOV-2001; 2001US-0334198P.

(CURA-) CURAGEN CORP.

Decristofaro MF, Padigaru M, Miller C, Tchernev V, Zhong H; Zhong M, Anderson D, Ballinger R, Gerlach V, Spytek KA, Rastelli L; Kekuda R, Guo X, Zernusen B, Andrew D, Mezes P, Patturajan M; Burgess CE, Eisen A, Wolenc A, Baumgartner J, Shinkets RA, Gusev V; Vermet CAM, Taupier RJ, Pena C, Shenoy S, Li L, Casman S, Boldog F; Fernandes B, Smithson G, Malyankar U, Tallon B, Liu X; WPI; 2003-058504/05.

P-PSDB; ABJ37909.

New polypeptides, designated as NOVX, useful for diagnosing and treating infections, neurological diseases, cancer, allergy, and bone, immunological, skin, renal, brain, muscle and autoimmune disorders.

Claim 9; Page 171; 672pp; English.

The invention relates to a novel isolated polypeptide, designated NOVX (NOV1 - 33), consisting of a mature form of one of 61 sequences, given in the specification, or its variant, where amino acid residue(s) in the variant differ from the mature form, provided that the variant differs in not more than 15 % of the amino acids from the sequence of the mature form. The NOVX polypeptides, nucleic acids encoding the polypeptides, and an antibody to the polypeptides, are useful for treating or preventing a NOVX-associated disorder in humans and for treating a syndrome associated with a human disease (NOVX-associated disorder). NOVX polypeptides and the encoding nucleic acids, are useful for determining the presence of or predisposition to a disease associated with altered levels of NOVX polypeptide and polynucleotide, by measuring the level of polypeptide expression or the amount of nucleic acid from a mammal and comparing it with another mammal not having or not predisposed to the disease. NOVX polypeptide is also useful for identifying an agent that binds to NOVX and a cell expressing NOVX is useful for identifying an agent that

modulates the expression or activity of NOVX. The antibodies and a polypeptide having 95 % sequence identity to NOVX polypeptide are useful for treating a pathological state in a mammal. The antibodies are also useful for determining the presence or amount of NOVX in a sample. NOVX polypeptides, polynucleotides and antibodies specific for the polypeptides are useful for treating or preventing disorders or syndromes including trauma, viral, bacterial, fungal, protozoal, and parasitic infections. They can also treat disorders such as e.g., Alzheimer's disease or a stroke. The NOVX encoding nucleic acids are useful for expressing the NOVX proteins, to detect NOVX mRNA, or a genetic lesion in a NOVX gene and to modulate NOVX activity. NOVX sequences are also useful for identifying a cell or tissue type in a biological sample, to amplify DNA sequences from very small biological samples such as tissues e.g. hair or skin or body fluids in forensic biology and as primers and probes for use in identifying and/or cloning NOVX homologues in other cell types. The NOVX proteins are useful as an immunogen to generate antibodies which are useful for diagnostically monitoring protein levels and modulating NOVX activity. Cells comprising NOVX nucleic acids are useful for producing non-human transgenic animals which are useful for studying the function and/or activity of NOVX protein and for identifying and/or evaluating modulators of NOVX protein activity. The NOVX nucleic acids can be used in gene therapy. This polynucleotide sequence represents a NOVX DNA sequence of the invention

SQ Sequence 2463 BP; 691 A; 599 C; 580 G; 593 T; 0 U; 0 Other;

Query Match 41.2%; Score 410; DB 7; Length 2463;
Best Local Similarity 100.0%; Pred. No. 1.5e-190;
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 450 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGGCAGAGCGATGAGTTC 509
Db 2009 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGGCAGAGCGATGAGTTC 2068

Qy 510 CTTAGTATCATGTCATGACAGATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 569
Db 2069 CTTAGTATCATGTCATGACAGATGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 2128

Qy 570 CCGCATGGTGTATTAATAATTCATACACAAAGAGAGATTAGACAGTAAAGAGTTTACAAG 629
Db 2129 CCGCATGGTGTATTAATAATTCATACACAAAGAGAGATTAGACAGTAAAGAGTTTACAAG 2188

Qy 630 AATAAATCTATATTTTGTGAAGGTAGTGGTATTATATCTGTAGATTTTCAGTAGTTTCT 699
Db 2189 AATAAATCTATATTTTGTGAAGGTAGTGGTATTATATCTGTAGATTTTCAGTAGTTTCT 2248

Qy 690 AAGTCTGTATTCGTTTGTAAACAATGGCAGGTTTACACGCTCTATGCAATTCACAAA 749
Db 2249 AAGTCTGTATTCGTTTGTAAACAATGGCAGGTTTACACGCTCTATGCAATTCACAAA 2308

Qy 750 AAGTTATAAGAAAACACTACATGTAATAATCTTGTATGAGTAAATAACTTGCATTTCTTTATA 809
Db 2309 AAGTTATAAGAAAACACTACATGTAATAATCTTGTATGAGTAAATAACTTGCATTTCTTTATA 2368

Qy 810 TGGNACGCATTTGGGTGTTTAAATAATTTATAACAGTTTATAAGAAAGA 859
Db 2369 TGGNACGCATTTGGGTGTTTAAATAATTTATAACAGTTTATAAGAAAGA 2418

RESULT 5
ABK81998
ID ABK81998 standard; DNA; 5339 BP.
AC ABK81998;
XX
XX 13-AUG-2002 (first entry)
XX DNA encoding mini-dystrophin protein deltaR2-R21.
XX Mini-dystrophin peptide; spectrin-like repeat domain; muscle disease;
KW Duchenne's muscular dystrophy; DMD; dystrophin; ds.
XX
XX Homo sapiens.
OS

```

OS Synthetic.
XX WO200229056-A2.
XX
XX
XX 11-APR-2002.
XX
XX 04-OCT-2001; 2001WO-US031126.
XX
XX 06-OCT-2000; 2000US-0238848P.
XX
XX (UNMI ) UNIV MICHIGAN.
XX
XX Chamberlain JS, Harper SQ;
XX
XX WPI; 2002-435334/46.
XX
XX
XX A composition for preparing therapeutic drugs, has a mini-dystrophin
XX peptide comprising a specific number of spectrin-like repeat domains, or
XX a nucleic acid sequence encoding the mini-dystrophin peptide.
XX
XX Example 6; Fig 13; 145pp; English.
XX
XX The invention describes a composition comprising a mini-dystrophin
XX peptide comprising a spectrin-like repeat domain, where the domain
XX comprises n spectrin-like repeats, and contains no more than n spectrin-
XX like repeats, where n is an even number between 4-24, or a nucleic acid
XX encoding a mini-dystrophin peptide. The mini-dystrophin peptide or the
XX polynucleotide encoding it is useful as a medicament, for preparing a
XX drug for therapeutic application and in the preparation of a composition
XX for treatment of muscle disease, e.g. Duchenne's muscular dystrophy
XX (DMD). This sequence represents a mini-dystrophin sequence of the
XX invention
XX
XX Sequence 5339 BP; 1638 A; 1191 C; 1187 G; 1323 T; 0 U; 0 Other;
XX
XX Query Match 41.2%; Score 410; DB 6; Length 5339;
XX Best Local Similarity 100.0%; Pred. No. 1.5e-190;
XX Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 450 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGGCAGAGCGATGGAGTC 509
XX |
XX Db 4491 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGGCAGAGCGATGGAGTC 4550
XX |
XX QY 510 CTTAGTATCATGTGATGACAGATGAAGAGGAGCGAATAAATGTTTTCACACTCCTGATT 569
XX |
XX Db 4551 CTTAGTATCATGTGATGACAGATGAAGAGGAGCGAATAAATGTTTTCACACTCCTGATT 4610
XX |
XX QY 570 CCGCATGGTTTTTATATATTCATACAAACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
XX |
XX Db 4611 CCGCATGGTTTTTATATATTCATACAAACAAAGAGGATTAGACAGTAAGAGTTTACAAG 4670
XX |
XX QY 630 AAATAAATCTATATTTTTGTGAAGGGTAGTGGTATTAATCTAGTAGATTTTCAGTAGTTTCT 689
XX |
XX Db 4671 AAATAAATCTATATTTTTGTGAAGGGTAGTGGTATTAATCTAGTAGATTTTCAGTAGTTTCT 4730
XX |
XX QY 690 AAGTCTGTTATGTTTTGTTTAAACAATGGCAGGTTTTACAGTCTATCGCAATTGTACAAA 749
XX |
XX Db 4731 AAGTCTGTTATGTTTTGTTTAAACAATGGCAGGTTTTACAGTCTATCGCAATTGTACAAA 4790
XX |
XX QY 750 AAGTTATAAGAAAACTACATGTAATAATCTTGATAGCTAAATAAAGTTCGCCATTTCTTTATA 809
XX |
XX Db 4791 AAGTTATAAGAAAACTACATGTAATAATCTTGATAGCTAAATAAAGTTCGCCATTTCTTTATA 4850
XX |
XX QY 810 TGGAACCCATTTTGGGTGTTTTAAAAAATTTATTAACAGTTTAAAGAAGA 859
XX |
XX Db 4851 TGGAACCCATTTTGGGTGTTTTAAAAAATTTATTAACAGTTTAAAGAAGA 4900
XX |
XX
XX RESULT 6
XX ABK81997
XX ID ABK81997 standard; DNA; 5417 BP.
XX
XX AC ABK81997;

```

Db 4929 TCGAAGCGCATTTTGGGTGTTTAAAAAATTTATAACAGTATATAAGAAAGA 4978

RESULT 7

ABK81999

ID ABK81999 standard; DNA; 5462 BP.

XX ABK81999;

XX AC

XX DT

XX 13-AUG-2002 (first entry)

XX DE

XX DNA encoding mini-dystrophin protein deltar2-R21-H3.

DE DE

XX Mini-dystrophin peptide; spectrin-like repeat domain; muscle dis

KW Mini-dystrophin peptide; spectrin-like repeat domain; muscle dis

KW Duchenne's muscular dystrophy; DMD; dystrophin; ds.

XX OS

XX Homo sapiens.

OS OS

XX Synthetic.

XX PN

XX W0200229056-A2.

XX PD

XX 11-APR-2002.

XX PF

XX 04-OCT-2001; 2001WO-US031126.

XX PR

XX 06-OCT-2000; 2000US-0238848P.

XX PA

XX (UNMI) UNIV MICHIGAN.

XX PI

XX Chamberlain JS, Harper SQ;

XX DR

XX WPI; 2002-435334/46.

XX PT

XX A composition for preparing therapeutic drugs, has a mini-dystro

PT peptide comprising a specific number of spectrin-like repeat dom

PT a nucleic acid sequence encoding the mini-dystrophin peptide.

XX PS

XX Disclosure; Fig 14; 145pp; English.

XX CC

XX The invention describes a composition comprising a mini-dystroph

CC peptide comprising a spectrin-like repeat domain, where the doma

CC comprises n spectrin-like repeats, and contains no more than n s

CC like repeats, where n is an even number between 4-24, or a nucle

CC encoding a mini-dystrophin peptide. The mini-dystrophin peptide

CC polynucleotide encoding it is useful as a medicament, for prepar

CC drug for therapeutic application and in the preparation of a com

CC for treatment of muscle disease, e.g. Duchenne's muscular dystro

CC (DMD). This sequence represents a mini-dystrophin sequence of th

XX invention

XX SQ

SQ Sequence 5462 BP; 1668 A; 1225 C; 1212 G; 1357 T; 0 U; 0 Other;

Query Match 41.2%; Score 410; DB 6; Length 5462;

Best Local Similarity 100.0%; Pred. No. 1.5e-190;

Matches 410; Conservative 0; Mismatches 0; Indels 0;

QY 450 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTGGCGCAGCGCATG

Db 4614 AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTGGCGCAGCGCATG

QY 510 CTTAGTATCAGTCATCAGCAGATGAGAAGGAGCAGAAATAATGTTTTACAACCTCC

Db 4674 CTTAGTATCAGTCATCAGCAGATGAGAAGGAGCAGAAATAATGTTTTACAACCTCC

QY 570 CCGCATGGTGTTTTAATAATTCATACAAAGAGGATTAGACAGTAAGAGTATT

Db 4734 CCGCATGGTGTTTTAATAATTCATACAAAGAGGATTAGACAGTAAGAGTATT

QY 630 AAATAAATCTAATTTTTGTGAAGGAGTAGTGTAATTA TACTCTAGATTT CAGTAGT

Db 4794 AAATAAATCTAATTTTTGTGAAGGAGTAGTGTAATTA TACTCTAGATTT CAGTAGT

Qy	590	AACTCTGTTATTCTTTTGTAAACAATGGCAGGTTTACACGCTCTATGCATTCGTACAAA	749			
Db	4854	AACTCTGTTATTCTTTTGTAAACAATGGCAGGTTTACACGCTCTATGCATTCGTACAAA	4913			
Qy	750	AACTTATAAGAAAACACTACATGTAATAATCTTTAGTAGCTAAATAACTTGGCCATTTCTTTTATA	809			
Db	4914	AACTTATAAGAAAACACTACATGTAATAATCTTTAGTAGCTAAATAACTTGGCCATTTCTTTTATA	4973			
Qy	810	TGGAAGCGCATTTTGGGTGTTTAAAAATTTTATAACAGTTTATAAGAAAGA	859			
Db	4974	TGGAAGCGCATTTTGGGTGTTTAAAAATTTTATAACAGTTTATAAGAAAGA	5023			
RESULT 8						
ABK82000						
ID	ABK82000-standard; DNA; 8689 BP.					
XX						
AC	ABK82000;					
XX						
DT	13-AUG-2002 (first entry)					
XX						
DE	DNA encoding mini-dystrophin protein deltaH2-R19.					
XX						
KW	Mini-dystrophin peptide; spectrin-like repeat domain; muscle disease;					
KW	Duchenne's muscular dystrophy; DMD; dystrophin; ds.					
XX						
OS	Homo sapiens.					
OS	Synthetic.					
XX						
FN	W0200229056-A2.					
XX						
PD	11-APR-2002.					
XX						
XX	04-OCT-2001; 2001WO-US031126.					
PF						
XX	06-OCT-2000; 2000US-0238848P.					
ER						
XX	(UNMI) UNIV MICHIGAN.					
PA						
XX	Chamberlain JS, Harper SQ;					
PI						
XX	WPI; 2002-435334/46.					
DR						
XX						
PT	A composition for preparing therapeutic drugs, has a mini-dystrophin					
PT	peptide comprising a specific number of spectrin-like repeat domains, or					
PT	a nucleic acid sequence encoding the mini-dystrophin peptide.					
XX						
PS	Disclosure; Fig 15; 145pp; English.					
XX						
CC	The invention describes a composition comprising a mini-dystrophin					
CC	peptide comprising a spectrin-like repeat domain, where the domain					
CC	comprises n spectrin-like repeats, and contains no more than n spectrin-					
CC	like repeats, where n is an even number between 4-24, or a nucleic acid					
CC	encoding a mini-dystrophin peptide. The mini-dystrophin peptide or the					
CC	polynucleotide encoding it is useful as a medicament, for preparing a					
CC	drug for therapeutic application and in the preparation of a composition					
CC	for treatment of muscle disease, e.g. Duchenne's muscular dystrophy					
CC	(DMD). This sequence represents a mini-dystrophin sequence of the					
XX						
SQ	Sequence 8689 BP; 2721 A; 1804 C; 1861 G; 2303 T; 0 U; 0 Other;					
Query March 41.2%; Score 410; DB 6; Length 8689;						
Best Local Similarity 100.0%; Pred. No. 1.5e-190;						
Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;						
Qy	450	AGGACACAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGGCAGAGCGATGGAGTC	509			
Db	5985	AGGACACAATGTAGGAAGTCTTTTCCATGGCAGATGATTTGGGCAGAGCGATGGAGTC	6044			
Qy	510	CTTAGTATCAGTCATGACAGATGAAGAGAGAGAGATAAATGTTTACAACCTCTGATT	569			
Db	6045	CTTAGTATCAGTCATGACAGATGAAGAGAGAGAGATAAATGTTTACAACCTCTGATT	6104			

QY 570 CCGCATGGTCTTTTATTAATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
 DB 6105 CCGCATGGTCTTTTATTAATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG 6164
 QY 630 AATAAATCTATATTTTGTGAAGGGTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 689
 DB 6165 AATAAATCTATATTTTGTGAAGGGTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 6224
 QY 690 AGCTCTGTATTTTGTGTAAACAATGGCAGGTTTACACGCTATATGCAATTTGTACAAA 749
 DB 6225 AGCTCTGTATTTTGTGTAAACAATGGCAGGTTTACACGCTATATGCAATTTGTACAAA 6284
 QY 750 AGCTTATAGAAACACTACATGTAATCTTGTAGCTAAATAACTTGCATTTCTTTATA 809
 DB 6285 AGCTTATAGAAACACTACATGTAATCTTGTAGCTAAATAACTTGCATTTCTTTATA 6344
 QY 810 TGAACGCAATTTTGGTGTGTTTAAAAATTTTAAACAGTTTAAAGAAAGA 859
 DB 6345 TGAACGCAATTTTGGTGTGTTTAAAAATTTTAAACAGTTTAAAGAAAGA 6394

RESULT 9
 ABK82002
 ID ABK82002 standard; DNA; 11443 BP.

AC ABK82002;
 DT 13-AUG-2002 (first entry)

XX DNA encoding mini-dystrophin protein deltaR9-R16.

XX Mini-dystrophin peptide; spectrin-like repeat domain; muscle disease;
 XX Duchenne's muscular dystrophy; DMD; dystrophin; ds.

XX Homo sapiens.
 OS Synthetic.

XX WO200229056-A2.

XX 11-APR-2002.

XX 04-OCT-2001; 2001WO-US031126.

XX 06-OCT-2000; 2000US-0238848P.

XX (UNMI) UNIV MICHIGAN.

XX Chamberlain JS, Harper SQ;

XX WPI; 2002-435334/46.

XX A composition for preparing therapeutic drugs, has a mini-dystrophin
 PT peptide comprising a specific number of spectrin-like repeat domains, or
 PT a nucleic acid sequence encoding the mini-dystrophin peptide.

XX Disclosure; Fig 17; 145pp; English.

XX The invention describes a composition comprising a mini-dystrophin
 CC peptide comprising a spectrin-like repeat domain, where the domain
 CC comprises n spectrin-like repeats, and contains no more than n spectrin-
 CC like repeats, where n is an even number between 4-24, or a nucleic acid
 CC encoding a mini-dystrophin peptide. The mini-dystrophin peptide or the
 CC polynucleotide encoding it is useful as a medicament, for preparing a
 CC drug for therapeutic application and in the preparation of a composition
 CC for treatment of muscle disease, e.g. Duchenne's muscular dystrophy
 CC (DMD). This sequence represents a mini-dystrophin sequence of the
 CC invention

XX Sequence 11443 BP; 3707 A; 2339 C; 2502 G; 2895 T; 0 U; 0 Other;

XX Query Match 41.2%; Score 410; DB 6; Length 11443;

XX Best Local Similarity 100.0%; Pred. No. 1.5e-190;

Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 450 AGGACCAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGGAGAGCGATGAGTC 509
 DB 8739 AGGACCAATGTAGGAAGTCTTTTCCACATGGCAGATGATTTGGGAGAGCGATGAGTC 8798
 QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGAAATAAATGTTTTTACAACCTCGATT 569
 DB 8799 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAGAAATAAATGTTTTTACAACCTCGATT 8858
 QY 570 CCCCATGGTCTTTTATTAATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG 629
 DB 8859 CCCCATGGTCTTTTATTAATTCATACAAAGAGGATTAGACAGTAAGAGTTTACAAG 8918
 QY 630 AATAAATCTATATTTTGTGAAGGGTAGTGGTATTATCTAGTAGATTTTTCAGTAGTTTCT 689
 DB 8919 AATAAATCTATATTTTGTGAAGGGTAGTGGTATTATCTAGTAGATTTTTCAGTAGTTTCT 8978
 QY 690 AAGTCTGTATTTTGTGTTAAACAATGGCAGGTTTACACGCTCTATGCAATTTGTACAAA 749
 DB 8979 AAGTCTGTATTTTGTGTTAAACAATGGCAGGTTTACACGCTCTATGCAATTTGTACAAA 9038
 QY 750 AAGTTATAAGAAACTACATGTAATAATCTTGTAGCTAAATAAATCTGCCATTTCTTTATA 809
 DB 9039 AAGTTATAAGAAACTACATGTAATAATCTTGTAGCTAAATAAATCTGCCATTTCTTTATA 9098
 QY 810 TGAACGCAATTTTGGTGTGTTTAAAAATTTTAAACAGTTTAAAGAAAGA 859
 DB 9099 TGAACGCAATTTTGGTGTGTTTAAAAATTTTAAACAGTTTAAAGAAAGA 9148

RESULT 10

AA90338
 ID AA90338 standard; cDNA; 12923 BP.

XX AA90338;

DT 29-MAR-1992 (first entry)

XX Sequence of human muscular dystrophy (MD) cDNA.

XX Dystrophin; muscular dystrophy; probe; antibody; diagnosis; prenatal;
 KW heterozygote; gene therapy; genetic screening; foetal screening; ss.

XX Homo sapiens.

XX Key Location/Qualifiers
 FT CDS 209..12923
 FT /*tag= a

XX WO8906286-A.

XX 13-JUL-1989.

XX 16-DEC-1988; 88WO-US004504.

XX 22-DEC-1987; 87US-00136618.

XX (CHIL-) CHILDRENS MED CENT.

XX Kunkel LM, Monaco A, Hoffman EP, Koenig M;

XX WPI; 1989-220587/30.

XX P-PSDB; AAP90373.

XX Muscular dystrophy gene - used for prepn. of probes, dystrophic
 PT polypeptide and antibodies for diagnosis and therapy of muscular
 PT dystrophy.

XX Disclosure; Fig 5; 68pp; English.

XX The inventors claim an MD probe comprising a purified ss NA SQ which
 CC hybridises to at least a part of the MD gene; pure dystrophin (DS)

CC polypeptide, purified NA encoding DS and antibodies (Ab) to DS. The
 CC probes are equal to or greater than 10b of one of 12 cDNA sequences
 CC deposited as ATCC 58666-57677. The MD gene is human, or a murine Dmd gene
 XX
 SQ Sequence 12923 BP; 4296 A; 2613 C; 2963 G; 3044 T; 0 U; 7 Other;
 Query Match 41.2%; Score 410; DB 1; Length 12923;
 Best Local Similarity 100.0%; Pred. No. 1.5e-190;
 Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 450 AGGACACATGTAGGAAGTCTTTTCCACATGCGAGATGTTGGCAGAGCGATGGAGTC 509
 DB 11253 AGGACACATGTAGGAAGTCTTTTCCACATGCGAGATGTTGGCAGAGCGATGGAGTC 11312
 QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAATAAATGTTTCAACTCTCTGATT 569
 DB 11313 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAATAAATGTTTCAACTCTCTGATT 11372
 QY 570 CCGCATGCTTTTATTAATATTCATACACAAAGAGGATTAGACAGTAAGATTTTACAAG 629
 DB 11373 CCGCATGCTTTTATTAATATTCATACACAAAGAGGATTAGACAGTAAGATTTTACAAG 11432
 QY 630 AAATAAATCTATATTTTGTGAAGGCTAGTGTATTATCTAGATTTTCAAGTAGTTTCT 689
 DB 11433 AAATAAATCTATATTTTGTGAAGGCTAGTGTATTATCTAGATTTTCAAGTAGTTTCT 11492
 QY 690 AAGTCTGTTATTTTGTGAAGGCTAGTGTATTATCTAGATTTTCAAGTAGTTTCT 749
 DB 11493 AAGTCTGTTATTTTGTGAAGGCTAGTGTATTATCTAGATTTTCAAGTAGTTTCT 11552
 QY 750 AAGTTATAGAAACATACATGTAATAATCTTGATAGCTAAATTAACCTTGTCTTTTATA 809
 DB 11553 AAGTTATAGAAACATACATGTAATAATCTTGATAGCTAAATTAACCTTGTCTTTTATA 11612
 QY 810 TGGAAACGCAATTTGGTGTGTTTAAAAATTTTAAACAGTTTATAAAGAAAGA 859
 DB 11613 TGGAAACGCAATTTGGTGTGTTTAAAAATTTTAAACAGTTTATAAAGAAAGA 11662

RESULT 11
 ABK81959 standard; DNA; 13957 BP.
 ID ABK81959
 XX
 AC ABK81959;
 DT 13-AUG-2002 (first entry)
 XX
 DE cDNA encoding human dystrophin.
 XX
 KW Mini-dystrophin peptide; spectrin-like repeat domain; muscle disease;
 KW Duchenne's muscular dystrophy; DMD; dystrophin; human; gene; ds.
 XX
 OS Homo sapiens.
 XX
 DN WO200229056-A2.
 XX
 PD 11-APR-2002.
 XX
 XX 04-OCT-2001; 2001WO-US031126.
 XX
 XX 06-OCT-2000; 2000US-0238848P.
 XX
 XX (UNMI) UNIV MICHIGAN.
 XX
 XX Chamberlain JS, Harper SQ;
 XX WPI; 2002-435334/46.
 XX
 XX A composition for preparing therapeutic drugs, has a mini-dystrophin
 PT peptide comprising a specific number of spectrin-like repeat domains, or
 PT a nucleic acid sequence encoding the mini-dystrophin peptide.
 XX
 XX Example 2; Fig 1; 145pp; English.
 PS

XX The invention describes a composition comprising a mini-dystrophin
 CC peptide comprising a spectrin-like repeat domain, where the domain
 CC comprises n spectrin-like repeats, and contains no more than n spectrin-
 CC like repeats, where n is an even number between 4-24, or a nucleic acid
 CC encoding a mini-dystrophin peptide. The mini-dystrophin peptide or the
 CC polynucleotide encoding it is useful as a medicament, for preparing a
 CC drug for therapeutic application and in the preparation of a composition
 CC for treatment of muscle disease, e.g. Duchenne's muscular dystrophy
 CC (DMD). This sequence represents a human dystrophin polynucleotide
 CC sequence used in the creation of the mini-dystrophin peptides of the
 CC invention
 XX
 SQ Sequence 13957 BP; 4602 A; 2781 C; 3122 G; 3452 T; 0 U; 0 Other;
 Query Match 41.2%; Score 410; DB 6; Length 13957;
 Best Local Similarity 100.0%; Pred. No. 1.5e-190;
 Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 450 AGGACACATGTAGGAAGTCTTTTCCACATGCGAGATGTTGGCAGAGCGATGGAGTC 509
 DB 11253 AGGACACATGTAGGAAGTCTTTTCCACATGCGAGATGTTGGCAGAGCGATGGAGTC 11312
 QY 510 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAATAAATGTTTCAACTCTCTGATT 569
 DB 11313 CTTAGTATCAGTCATGACAGATGAAGAAGGAGCAATAAATGTTTCAACTCTCTGATT 11372
 QY 570 CCGCATGCTTTTATTAATATTCATACACAAAGAGGATTAGACAGTAAGATTTTACAAG 629
 DB 11373 CCGCATGCTTTTATTAATATTCATACACAAAGAGGATTAGACAGTAAGATTTTACAAG 11432
 QY 630 AAATAAATCTATATTTTGTGAAGGCTAGTGTATTATCTAGATTTTCAAGTAGTTTCT 689
 DB 11433 AAATAAATCTATATTTTGTGAAGGCTAGTGTATTATCTAGATTTTCAAGTAGTTTCT 11492
 QY 690 AAGTCTGTTATTTTGTGAAGGCTAGTGTATTATCTAGATTTTCAAGTAGTTTCT 749
 DB 11493 AAGTCTGTTATTTTGTGAAGGCTAGTGTATTATCTAGATTTTCAAGTAGTTTCT 11552
 QY 750 AAGTTATAGAAACATACATGTAATAATCTTGATAGCTAAATTAACCTTGTCTTTTATA 809
 DB 11553 AAGTTATAGAAACATACATGTAATAATCTTGATAGCTAAATTAACCTTGTCTTTTATA 11612
 QY 810 TGGAAACGCAATTTGGTGTGTTTAAAAATTTTAAACAGTTTATAAAGAAAGA 859
 DB 11613 TGGAAACGCAATTTGGTGTGTTTAAAAATTTTAAACAGTTTATAAAGAAAGA 11662

RESULT 12
 ABT10904 standard; cDNA; 13957 BP.
 ID ABT10904
 XX
 AC ABT10904;
 DT 04-DEC-2002 (first entry)
 XX
 DE Human breast cancer associated coding sequence SEQ ID NO: 1038.
 XX
 KW Human; breast specific gene; breast cancer; differential expression;
 KW cytostatic; gene therapy; gene; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200259271-A2.
 XX
 PD 01-AUG-2002.
 XX
 XX 25-JAN-2002; 2002WO-US002176.
 XX
 XX 25-JAN-2001; 2001US-0263757P.
 PR 25-APR-2001; 2001US-0286090P.
 PR 23-MAY-2001; 2001US-0292517P.
 XX

PA (GENE-) GENE LOGIC INC.
 XX Orr MS, Nation M, Diggins JC, Zeng W;
 XX WPI; 2002-674803/72.
 XX Diagnosing breast cancer in a patient comprises detecting the level of
 PT gene expression in cell or tissue samples, where a differential gene
 PT expression is indicative of breast cancer.
 XX
 XX Claim 1; SEQ ID NO 1038; 260pp + Sequence Listing; English.
 PS
 XX The present invention relates to methods of diagnosing breast cancer in a
 CC patient, which comprise detecting the level of expression in a tissue
 CC sample of two or more genes selected from those shown in AB09867-
 CC AB1112, where a differential expression of the genes indicates breast
 CC cancer. The methods are useful in diagnosing, treating, detecting the
 CC progression, and in monitoring treatment of breast cancer in patients.
 CC The methods are also useful as a screening tool for agents that modulate
 CC the onset or progression of breast cancer. The breast cancer genes may be
 CC used as diagnostic markers for breast cancer. The prediction or identification of the
 CC malignant state of breast tissue, for confirming the type and progression
 CC of cancer, and for drug screening and assays. The present sequence is a
 CC coding sequence of the invention. Note: The sequence data for this patent
 CC did not form part of the printed specification, but was obtained in
 CC electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 XX Sequence 13957 BP; 4602 A; 2781 C; 3122 G; 3452 T; 0 U; 0 Other;
 SQ
 Query Match 41.2%; Score 410; DB 6; Length 13957;
 Best Local Similarity 100.0%; Pred. No. 1.5e-190;
 Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 450 AGGACACATGTAGGAGTCTTTTCCACATGGCAGATGTTGGGACAGCGATGAGTC 509
 DB 11253 AGGACACATGTAGGAGTCTTTTCCACATGGCAGATGTTGGGACAGCGATGAGTC 11312
 QY 510 CTTAGTATCATGATCAGATGAAGAGGAGCAGATGAATGTTTACAACTCTCTGATT 569
 DB 11313 CTTAGTATCATGATCAGATGAAGAGGAGCAGATGAATGTTTACAACTCTCTGATT 11372
 QY 570 CCGCATGTTTATTAATATTTTGAAGGGTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 629
 DB 11373 CCGCATGTTTATTAATATTTTGAAGGGTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 11432
 QY 630 AAATAAATCTATATTTTGTGAAGGGTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 689
 DB 11433 AAATAAATCTATATTTTGTGAAGGGTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 11492
 QY 690 AGTCTGTTATTTGTTTAAACATGGCAGGTTTACAGCTCTATGCAATTGTACAAA 749
 DB 11493 AGTCTGTTATTTGTTTAAACATGGCAGGTTTACAGCTCTATGCAATTGTACAAA 11552
 QY 750 AAGTTATAAGAAACTACATGTAATAATCTTGTAGTCTAAATACTTGCATTTCTTTATA 809
 DB 11553 AAGTTATAAGAAACTACATGTAATAATCTTGTAGTCTAAATACTTGCATTTCTTTATA 11612
 QY 810 TGAACCGCATTTGGGTTGTTTAAATAATTTTAAACAGTTATAAGAAAGA 859
 DB 11613 TGAACCGCATTTGGGTTGTTTAAATAATTTTAAACAGTTATAAGAAAGA 11662
 RESULT 13
 ABN95786
 ID ABN95786 standard; DNA; 13957 BP.
 XX
 AC ABN95786;
 XX
 DT 13-AUG-2002 (first entry)
 XX
 DE Gene #2284 used to diagnose liver cancer.
 XX

KW Gene; liver cancer; ds; hepatocellular carcinoma; hepatotropic;
 KW metastatic liver tumour; cytostatic; expression profile; disease state;
 KW disease progression; drug toxicity; drug efficacy; drug metabolism.
 XX Homo sapiens.
 XX WO200229103-A2.
 XX 11-APR-2002.
 XX 02-OCT-2001; 2001WO-US030589.
 XX 02-OCT-2000; 2000US-0237054P.
 XX (GENE-) GENE LOGIC INC.
 XX Horne D, Alvares C, Peres-Da-Silva S, Vockley JG;
 XX WPI; 2002-426119/45.
 XX Diagnosing and detecting the progression of liver cancer, hepatocellular
 PT carcinoma or metastatic liver tumor in a patient, involves detecting the
 PT level of expression of two or more genes in a liver tissue sample.
 XX
 PS Claim 1; SEQ ID NO 2284; 298pp; English.
 XX The invention relates to a novel method for diagnosing and detecting the
 CC progression of liver cancer, hepatocellular carcinoma or metastatic liver
 CC tumor in a patient, and differentiating metastatic liver cancer from
 CC hepatocellular carcinoma in a patient, involving detecting the level of
 CC expression of two or more genes represented in ABN93503-ABN97455 in a
 CC tissue sample. The method of the invention has hepatotropic, and
 CC cytostatic activity. The method is useful for diagnosing and detecting
 CC the progression of liver cancer, hepatocellular carcinoma and metastatic
 CC liver carcinoma in a patient. The method is useful for identifying
 CC expression profiles which serve as useful diagnostic markers as well as
 CC markers that can be used to monitor disease states, disease progression,
 CC drug toxicity, drug efficacy and drug metabolism. Note: The sequence data
 CC for this patent did not form part of the printed specification, but was
 CC obtained in electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 XX Sequence 13957 BP; 4602 A; 2781 C; 3122 G; 3452 T; 0 U; 0 Other;
 SQ
 Query Match 41.2%; Score 410; DB 6; Length 13957;
 Best Local Similarity 100.0%; Pred. No. 1.5e-190;
 Matches 410; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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 QY 510 CTTAGTATCATGATCAGATGAAGAGGAGCAGATGAATGTTTACAACTCTCTGATT 569
 DB 11313 CTTAGTATCATGATCAGATGAAGAGGAGCAGATGAATGTTTACAACTCTCTGATT 11372
 QY 570 CCGCATGTTTATTAATATTTTGAAGGGTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 629
 DB 11373 CCGCATGTTTATTAATATTTTGAAGGGTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 11432
 QY 630 AAATAAATCTATATTTTGTGAAGGGTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 689
 DB 11433 AAATAAATCTATATTTTGTGAAGGGTAGTGGTATTATATCTAGATTTTCAGTAGTTTCT 11492
 QY 690 AGTCTGTTATTTGTTTAAACATGGCAGGTTTACAGCTCTATGCAATTGTACAAA 749
 DB 11493 AGTCTGTTATTTGTTTAAACATGGCAGGTTTACAGCTCTATGCAATTGTACAAA 11552
 QY 750 AAGTTATAAGAAACTACATGTAATAATCTTGTAGTCTAAATACTTGCATTTCTTTATA 809
 DB 11553 AAGTTATAAGAAACTACATGTAATAATCTTGTAGTCTAAATACTTGCATTTCTTTATA 11612
 QY 810 TGAACCGCATTTGGGTTGTTTAAATAATTTTAAACAGTTATAAGAAAGA 859

